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Audrey Chanet, Dragan Milenkovic, Sylvain Claude, Jeanette A. M. Maier, Muhammad Kamran Khan, et al.. Flavanone metabolites decrease monocyte adhesion to TNF-alpha-activated endothelial cells by modulating expression of atherosclerosis-related genes. *British Journal of Nutrition*, 2013, 110 (4), pp.587-598. hal-02642629

HAL Id: hal-02642629

<https://hal.inrae.fr/hal-02642629>

Submitted on 28 May 2020

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Flavanone metabolites decrease monocyte adhesion to TNF- α -activated endothelial cells by modulating expression of atherosclerosis-related genes

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(Submitted 11 April 2012 – Final revision received 8 November 2012 – Accepted 8 November 2012)

Abstract

Flavanones are found specifically and abundantly in citrus fruits. Their beneficial effect on vascular function is well documented. However, little is known about their cellular and molecular mechanisms of action in vascular cells. The goal of the present study was to identify the impact of flavanone metabolites on endothelial cells and decipher the underlying molecular mechanisms of action. We investigated the impact of naringenin and hesperetin metabolites at 0.5, 2 and 10 μM on monocyte adhesion to TNF- α -activated human umbilical vein endothelial cells (HUVEC) and on gene expression. Except hesperetin-7-glucuronide and naringenin-7-glucuronide (N7G), when present at 2 μM , flavanone metabolites (hesperetin-3'-sulphate, hesperetin-3'-glucuronide and naringenin-4'-glucuronide (N4'G)) significantly attenuated monocyte adhesion to TNF- α -activated HUVEC. Exposure of both monocytes and HUVEC to N4'G and N7G at 2 μM resulted in a higher inhibitory effect on monocyte adhesion. Gene expression analysis, using TaqMan Low-Density Array, revealed that flavanone metabolites modulated the expression of genes involved in atherogenesis, such as those involved in inflammation, cell adhesion and cytoskeletal organisation. In conclusion, physiologically relevant concentrations of flavanone metabolites reduce monocyte adhesion to TNF- α -stimulated endothelial cells by affecting the expression of related genes. This provides a potential explanation for the vasculoprotective effects of flavanones.

Key words: Flavanones; Metabolites; Monocyte adhesion; Gene expression; Endothelial cells

Atherosclerosis, a multifactorial disease of the large arteries and the leading cause of CVD, is characterised by inflammatory and oxidative stress involving chronic dysfunction of the endothelium. This altered function of endothelial cells is associated with both an imbalance in the production of vasodilators/vasoconstrictors and an increase in the expression of pro-inflammatory, chemotactic and adhesion molecules^(1,2). These changes facilitate the adhesion of circulating immune cells to the activated endothelium, and their subsequent migration into the arterial wall constitutes early events in the

atherogenic process. Once in the intima, monocytes differentiate into activated macrophages, which take up oxidised lipoproteins, leading to the formation of foam cells⁽³⁾.

A positive association between a high consumption of fruits and vegetables and a reduced risk of CVD has been described^(4,5). Among the protective bioactive compounds found in fruits and vegetables, polyphenols seem to be of particular interest. Indeed, epidemiological studies have established an inverse relationship between the intake of flavonoids, a subclass of polyphenols, and mortality from CVD⁽⁶⁾. Animal

Abbreviations: ACTR, actin-related protein; CCL2, chemokine (C-C motif) ligand 2; EZR, ezrin; H3'G, hesperetin-3'-glucuronide; H3'S, hesperetin-3'-sulphate; H7G, hesperetin-7-glucuronide; H, hesperetin; HUVEC, human umbilical vein endothelial cell; ICAM-1, intercellular adhesion molecule 1; N4'G, naringenin-4'-glucuronide; N7G, naringenin-7-glucuronide; RAC, ras-related C3 botulinum toxin substrate 1; TLDA, TaqMan Low-Density Array; VCAM-1, vascular cell adhesion molecule 1.

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studies have demonstrated that polyphenols may exert anti-atherogenic effects through anti-oxidative and anti-inflammatory actions that may prevent endothelial dysfunction⁽⁷⁾.

Flavanones are a subclass of flavonoids found specifically and abundantly in citrus fruits. Recently, a prospective study demonstrated a positive association between the dietary intake of flavanones and flavanone-rich foods and reduced risk of CHD and stroke^(8,9). In addition, a recent clinical trial established that flavanone supplementation in healthy, middle-aged, moderately overweight men improved postprandial microvascular endothelial reactivity and lowered diastolic blood pressure after 1 month⁽¹⁰⁾. In animal studies, flavanone supplementation has been reported to exert anti-atherogenic properties in various animal models of diet-induced atherosclerosis^(11–13). Such effects have been associated with reduced expression of vascular cell adhesion molecule 1 (VCAM-1) and monocyte chemoattractant protein-1 in the aorta⁽¹¹⁾. We have also shown, in a recent study, that the anti-atherogenic effect of flavanones was associated with decreased concentrations of plasma biomarkers of endothelial dysfunction and modulation of expression of genes involved in adhesion and transendothelial migration⁽¹⁵⁾. Furthermore, the few studies that have investigated the effects of flavanones on endothelial cell activity *in vitro* have reported changes in the levels of cell adhesion molecules and reduced adhesion of monocytes to endothelial cells^(14–16). Altogether, these data suggest a beneficial role of flavanone consumption on vascular function.

Cell culture-based studies are important to identify the cellular and molecular mechanisms involved in flavanones' action regarding CVD prevention. However, to date, most of the studies performed have used supraphysiological concentrations of flavanones (aglycone or glycoside forms), which are not present in the circulation after citrus fruit consumption^(14,15). Indeed, after ingestion, flavanone glycosides naturally present in citrus fruits are first hydrolysed in the distal part of the intestine by the microflora to release aglycones, which are then absorbed. In the intestinal cells, aglycone forms are subjected to phase II biotransformations (glucuronidation, sulphation and methylation), processes which continue in the liver⁽¹⁷⁾. Consequently, flavanones present in plasma are not native or aglycone forms, but almost exclusively conjugated metabolites. Such metabolites were identified in human subjects as being hesperetin-7-glucuronide (H7G), hesperetin-3'-sulphate (H3'S), hesperetin-3'-glucuronide (H3'G), naringenin-4'-glucuronide (N4'G) and naringenin-7-glucuronide (N7G) after consumption of orange and grapefruit juice^(18–20). Whether these metabolites at physiologically relevant concentrations have biological potencies remain unknown.

The aim of the present study was to investigate the effect of flavanone metabolites on endothelial cell function, particularly on monocyte adhesion on TNF- α -activated endothelial cells. Furthermore, to decipher the underlying molecular mechanisms, the impact of flavanone metabolites on the expression profile of genes and proteins involved in atherosclerosis development have been analysed.

Material and methods

Materials and chemicals

Naringenin and hesperetin were purchased from Extrasynthese. Flavanone glucuronides (H7G, H3'G and N7G) and H3'S were chemically synthesised⁽²⁰⁾. Chemical structures of naringenin and hesperetin and their main conjugated forms are presented in Supplemental Fig. S1 (available online). Solutions of naringenin, hesperetin as well as of their conjugates were obtained by dissolving them in 50% ethanol in bidistilled water. TNF- α was obtained from R&D Systems and was dissolved in 0.1% PBS/bovine serum albumin (Pan Biotech) to obtain a stock solution of 100 μ g/ml.

Cell culture

Primary human umbilical vein endothelial cells (HUVEC; Lonza) were used at passage 5 and were cultured in a phenol-red-free endothelial growth medium supplemented with 2% fetal bovine serum, 0.4% fibroblast growth factor, 0.1% vascular endothelial growth factor, 0.1% heparin, 0.1% insulin-like growth factor, 0.1% ascorbic acid, 0.1% epidermal growth factor and 0.04% hydrocortisone (all from Lonza). Experiments were performed in twenty-four-well plates (Becton Dickinson).

A human monocytic cell line (U937) (American Type Culture Collection) was cultured in Roswell Park Memorial Institute medium (Pan Biotech) supplemented with 2% fetal bovine serum (Sigma). Both cultures were maintained at 37°C and 5% CO₂.

Monocyte adhesion assay

HUVEC were allowed to proliferate until they reached 60–70% confluence. The medium was then replaced to expose the cells for 24 h to an experimental medium containing solvent alone (ethanol 0.5%, control wells and vehicle), flavanones or flavanone metabolites (0.5, 2 or 10 μ M). At the end of the 24 h period, the confluent monolayer was stimulated for 4 h with 0.1 ng/ml TNF- α or PBS/bovine serum albumin (0.01%, negative control). Following TNF- α stimulation, 50 μ l of a 5×10^6 /ml U937 cell suspension were added to each well and cells were further incubated for 1 h. Non-adhering U937 cells were rinsed away using $1 \times$ PBS and the wells were fixed with 0.5% crystal violet in ethanol (Sigma). The number of attached U937 cells was counted for each well in three random microscopic fields defined by an eyepiece⁽²¹⁾. Triplicates for each condition were performed in three independent experiments.

In a second set of experiments, both HUVEC and U937 cells were pre-treated separately for 24 h with flavanones or flavanone metabolites (2 μ M). After 24 h, 50 μ l of a suspension of 5×10^6 /ml U937 cells, pre-exposed to flavanones or their metabolites, were added to the wells containing HUVEC pre-exposed to the same compounds and previously treated for 4 h with TNF- α . The two cell types were co-incubated for 1 h and the adhesion assay performed as described earlier.

The experiments were repeated three times in triplicate. The experimental approach is schematically represented in Fig. 1.

RNA extraction and complementary DNA synthesis

To collect RNA, sub-confluent HUVEC were exposed for 24 h to a medium containing solvent alone (ethanol 0.5%, control wells), flavanones or flavanone metabolites. The monolayer was stimulated for 5 h with TNF- α (0.1 ng/ml) in order to expose the cells to TNF- α for the same time period as in the adhesion assay, where the HUVEC were activated with TNF- α for 4 h prior to co-incubation with monocytes for a further 1 h. Following this step, the cells were rinsed twice with cold PBS and lysed (Qiagen). Samples were kept at -80°C before RNA extraction.

Total RNA was extracted using QIAGEN® RNeasy® Kits (Qiagen), according to the manufacturer's protocol. The quality of total RNA was monitored in 1% agarose gel by electrophoresis and RNA quantity was checked on a NanoDrop ND-1000 spectrophotometer (Thermo Scientific). RT of total RNA was performed using a High Capacity cDNA RT kit (Applied Biosystems), each reaction mixture containing 1 μ g RNA, 1 \times RT buffer, 4 mM-deoxyribonucleoside triphosphates (dNTP) mix, 1 \times random primers, 50U RT and 20U RNase inhibitor in a total volume of 20 μ l. The RT reactions were run under the following conditions: 25°C for 10 min, 37°C for 120 min and 85°C for 5 s.

Real-time quantitative PCR

Impact of flavanones and their metabolites on the expression of genes in HUVEC were performed using the TaqMan Low-Density Array (TLDA) (Applied Biosystems). The TLDA contained primers for ninety-six genes of interest. Genes were chosen according to their role in atherosclerosis development from published articles (Table S1, available online).

A measure of 200 ng (2 μ l) of complementary DNA was used in each sample and 98 μ l of nuclease-free water together with 100 μ l 2 \times TaqMan Universal PCR Master Mix (Applied Biosystems) were added for the quantitative real-time PCR measurements. This mixture was divided equally over two sample-loading ports of the TLDA. The arrays were centrifuged once (1 min, 1300 rpm at room temperature) to equally distribute the sample over the wells. Subsequently, the card was sealed to prevent an exchange between the wells. Quantitative PCR amplification was performed using an Applied Biosystems Prism 7900HT system (Applied Biosystems), with the following thermal cycler conditions: 2 min at 50°C and 10 min at 94.5°C, followed by forty cycles of 30 s at 97°C and 30 s at 59.7°C. Raw data were analysed using Sequence Detection System Software v2.4 (Applied Biosystems). Analyses were performed in triplicate.

Unsupervised modelling

The clustering analysis was performed on mRNA expression data. Unsupervised modelling was subsequently performed using hierarchical clustering, with the Euclidean distance for calculating the similarity between genes and the Ward's distance for the similarity between conditions using average linkage. Permutmatrix version 1.9.3 (<http://www.lirmm.fr/~caraux/PermutMatrix>) was used for hierarchical clustering⁽²²⁾.

Western blot analysis

For Western blot analysis of proteins, the same experimental conditions as for RNA extraction were used, i.e. the subconfluent HUVEC were exposed for 24 h to a medium containing solvent alone (ethanol 0.5%, control wells), flavanones or flavanone metabolites. The monolayer was stimulated for 5 h with TNF- α (0.1 ng/ml) in order to expose the cell to TNF- α for the same time period as in adhesion assay, where the HUVEC were activated with TNF- α for 4 h prior

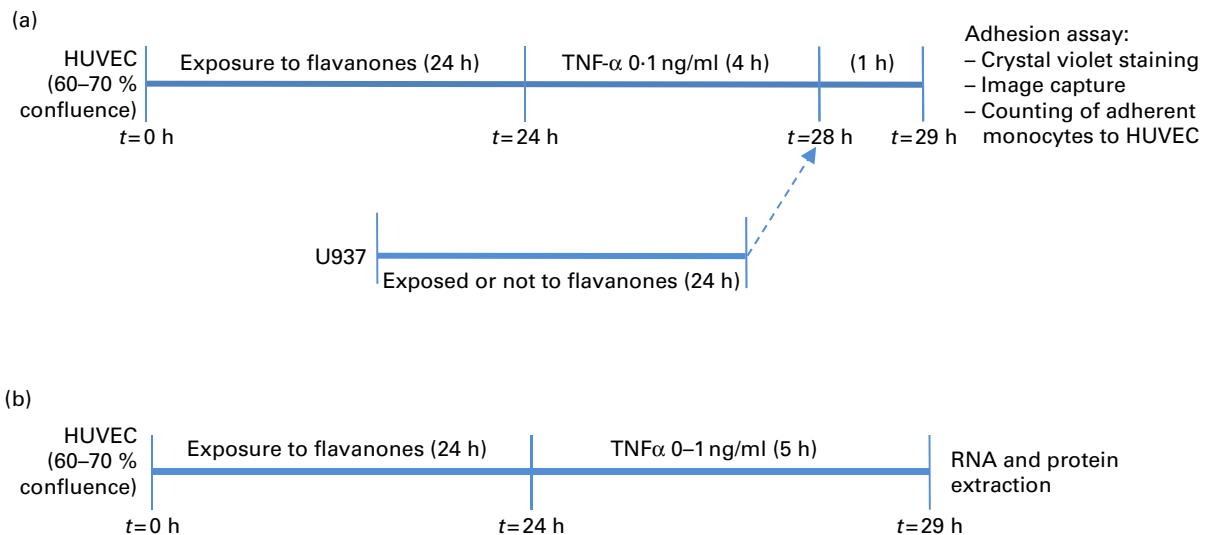


Fig. 1. Schematic representation of *in vitro* experimental approach. (a) Experimental condition for monocyte adhesion assay. (b) Experimental condition for RNA and protein extraction. HUVEC, human umbilical vein endothelial cells. (A colour version of this figure can be found online at <http://www.journals.cambridge.org/bjn>)

to co-incubation with monocytes for further 1 h. Following this step, the cells were rinsed twice with cold PBS and the total proteins were extracted using lysis buffer containing 150 mM-NaCl, 10 mM-Tris-HCl, 1 mM-ethylene glycol tetraacetic acid, 1 mM-EDTA, 100 mM-sodium fluoride, 4 mM-sodium pyrophosphate, 2 mM-sodium orthovanadate, 1% Triton X100 and 0.5% IGEPAL CA-630 (all were obtained from Sigma).

After washing with PBS, the cells were lysed using the lysis buffer supplemented with 1% protease inhibitor cocktail (Sigma) and the lysates were frozen at -80°C overnight after centrifugation at 14 000 rpm for 30 min at 4°C . Protein concentrations were determined using the BCA Protein Assay Kit (Interchim). A measure of $10\ \mu\text{g}$ of total proteins was electrophoretically separated on 8% polyacrylamide gels and transferred for 1 h on to nitrocellulose membranes Genetex (Euromedex). The membranes were then blocked for 1 h with 5% bovine serum albumin and Tris-buffered saline/0.1% Tween 20. After washing with Tris-buffered saline/0.1% Tween 20, the membranes were incubated with primary antibodies against intercellular adhesion molecule 1 (ICAM-1) (1/2000), VCAM-1 (1/2000), E-Selectin (1/2000), vinculin (1/2000) and β -actin (1/10 000); all the reagents were obtained from Genetex (Euromedex). The membranes were then incubated with secondary fluorescent goat anti-rabbit and goat anti-mouse IgG (H + L) antibodies (1/2000) from Genetex (Euromedex). Proteins were visualised by the Odyssey Li-Cor detection system (LI-COR) and the density of the signals was quantified using Odyssey software (LI-COR). Samples were normalised to the housekeeping protein β -actin. The expression of β -actin was used as the control for protein loading.

Statistical analyses

Adhesion assay. Data were analysed by one-way ANOVA, after Bartlett's test of homogeneity of variance and the Kolmogorov-Smirnov test for Gaussian distribution, followed by Dunnett's *post hoc* test. A value of $P < 0.05$ was considered significant. The software used was GraphPad InStat version 3.06 (GraphPad Software, Inc.).

Gene expression analysis. Relative gene expression levels of each gene were calculated by the comparative C_t method and the results were normalised to actin expression for each sample. Statistical analyses of the normalised gene expression data were performed using Dataassist software (Applied Biosystems; <http://products.appliedbiosystems.com/ab/en/US/adirect/ab?>, <http://cmd=catNavigate2&catID=606580>). The software utilises the comparative C_T ($\Delta\Delta C_T$) method to calculate quantitative relative gene expression across a large number of genes and samples. The software uses a refined Grubbs' outlier test to remove the outliers among technical replicates, provides a metric to measure control gene stability based on the geNorm algorithm to assist with endogenous control selection and allows the use of single or multiple control genes for data normalisation. $P < 0.05$ was taken to imply statistical significance.

Western blot. All experiments were run in three different cell preparations. Results are expressed as means and standard deviations. Statistical analyses were conducted using GraphPad InStat version 3.00 for Windows (GraphPad Software, Inc.). Significance was determined by Student's *t* test. $P < 0.05$ was considered statistically significant.

Results

Impact of pre-exposure of endothelial cells to flavanone metabolites on monocyte adhesion

Flavanones and their conjugated metabolites at a concentration of $10\ \mu\text{M}$ did not significantly influence cell viability compared with flavanone-free control solution (Supplementary data 1, available online).

As expected, a small number of monocytes adhered to unstimulated HUVEC and treatment with TNF- α significantly increased cell adhesion (280%, $P < 0.05$) compared with unstimulated control cells (Fig. 2). Pre-exposure of HUVEC to naringenin and N4'G (2 and $10\ \mu\text{M}$) for 24 h significantly reduced monocyte adherence to TNF- α -activated endothelial cells (-20%) (Fig. 2). A reduction in cell adhesion was also observed with N7G, but only at $10\ \mu\text{M}$ concentration (-24%). Pre-exposure of HUVEC to hesperetin (H), H3'G and H3'S also significantly reduced monocyte adhesion to activated endothelial cells at $2\ \mu\text{M}$ concentration (-23 , -21 and -14% , respectively), but this effect was no longer observed at a concentration of $10\ \mu\text{M}$. Finally, exposure to H7G did not produce any change in cell adhesion.

Impact of pre-exposure of both endothelial cells and monocytes to flavanone metabolites on cell adhesion

When both HUVEC and U937 cells were pre-exposed for 24 h with N4'G and N7G ($2\ \mu\text{M}$), the magnitude of the inhibition of cell adhesion was higher compared with the respective conditions with non-exposed monocytes (Fig. 3). For the two aglycone forms as well as for hesperetin metabolites, no differences were observed as compared with non-pretreated U937 cells.

Impact of flavanones and their metabolites on gene expression in TNF- α -stimulated endothelial cells

To evaluate the impact of flavanones and their metabolites on the expression of targeted genes, quantitative PCR was performed using TLDA. Out of ninety-six genes evaluated, eleven were identified as not expressed. Analysis of obtained quantitative PCR raw data by Dataassist software (Applied Biosystems) revealed that the expression of fourteen genes was significantly modified by TNF- α . Among these fourteen genes, expression of thirteen was up-regulated and the expression of one gene was down-regulated. Genes modulated by TNF- α are implicated in different cellular processes, such as cell adhesion (ICAM-1 and CDH5 (cadherin 5, type 2 - vascular endothelium)), inflammation and chemotaxis (N-FKB1 and chemokine (C-C motif) ligand 2

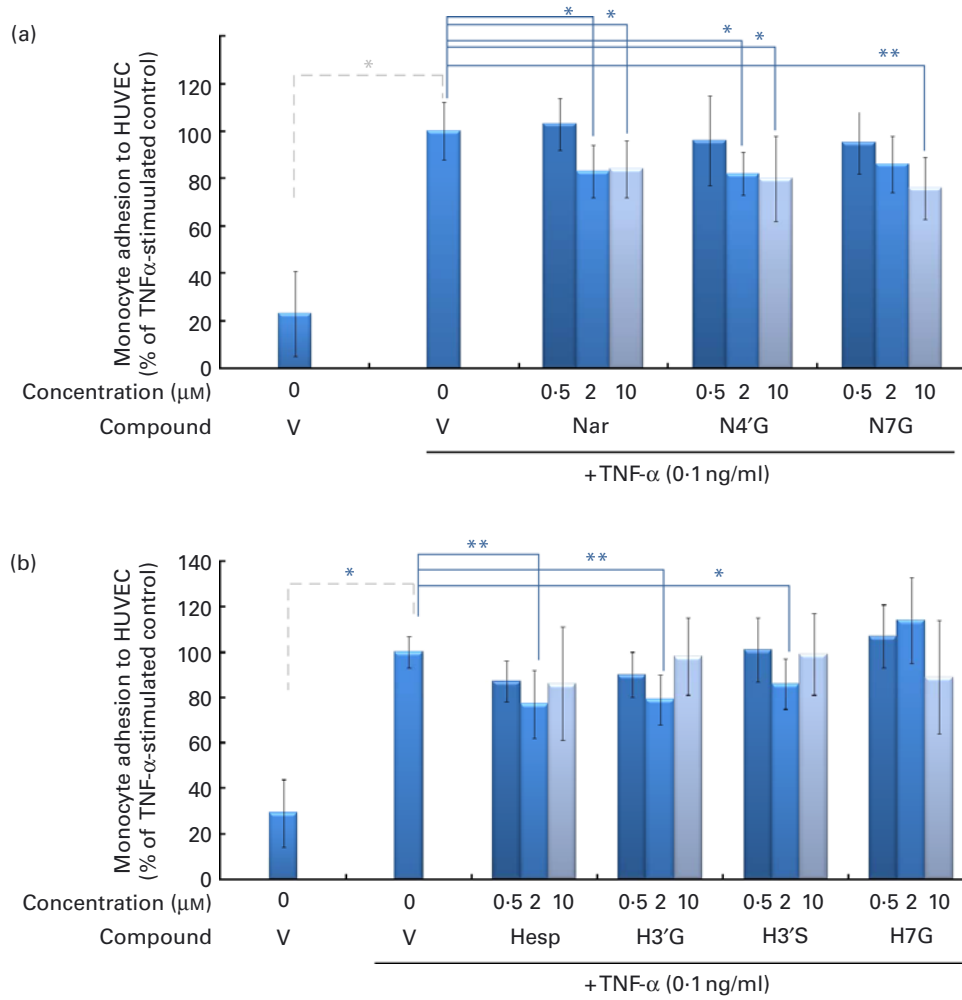


Fig. 2. Effect of incubation of endothelial cells with flavanones on monocyte adhesion. Human umbilical vein endothelial cells (HUVEC) were stimulated with TNF- α for 4 h after culture with or without (a) naringenin (Nar) and its conjugates (naringenin-4'-glucuronide (N4'G) and naringenin-7-glucuronide (N7G)) or (b) hesperetin (Hesp) and its conjugates (hesperetin-3'-glucuronide (H3'G); hesperetin-3'-sulphate (H3'S) and hesperetin-7-glucuronide (H7G)) for 24 h. Monocytes were co-incubated for 1 h and their adhesion to HUVEC counted. Monocyte adhesion is presented as a percentage of U937 to HUVEC. Values are means of three independent experiments, with standard deviations represented by vertical bars. Mean values were significantly different (by one-way ANOVA followed by Dunett's *post hoc* test): * $P < 0.05$, ** $P < 0.01$. (A colour version of this figure can be found online at <http://www.journals.cambridge.org/bjn>)

(CCL2)) or cytoskeleton organisation (actin-related protein (ACTR)2, ezrin (EZR) and RAC-1) (Fig. 4).

This gene expression study also revealed that the 24 h exposure of HUVEC to flavanones, aglycones or their metabolites at a physiologically relevant concentration (2 μM) before activation with TNF- α significantly modulated the expression of genes. H3'S modulated the expression of the highest number of genes, as twenty genes have been identified as differentially expressed after 24 h pre-exposition to this metabolite; H, H3'G, H7G and H3'S modulated the expression of fourteen, seven, six and twenty genes, respectively (Table S2, available online). Naringenin and its metabolites, N4'G and N7G, modulated the expression of three, nine and four genes, respectively (Table S2, available online). Taken as a whole, modulated genes are implicated in different cellular functions; for example, CD99 molecule (*CD99*), catenin (cadherin-associated protein), beta 1, 88 kDa (*CTNBN1*),

kelch-like ECH-associated protein 1 (*KEAP1*), moesin (*MSN*), integrin, beta 1 (*ITGB1*) and von Willebrand factor (*VWF*) are involved in cell-cell adhesion; *ACTR2*, *ACTR3*, platelet-derived growth factor receptor, alpha polypeptide (*PDGFRA*), p21 protein (Cdc42/Rac)-activated kinase 1 (*PAK1*), p21 protein (Cdc42/Rac)-activated kinase 4 (*PAK4*), *EZR* and protein phosphatase 1, regulatory subunit 12B (*PPP1R12B*) in actin cytoskeleton organisation; ras homolog family member A (*RHOA*) and cyclin-dependent kinase inhibitor 1 (*CDKN1A*) in Rho signalling transduction; and *CCL2*, *NFKB1* and mitogen-activated protein kinase 14 (*MAPK14*) in chemotaxis. As presented earlier, some of the cellular processes affected by flavanone metabolites have also been identified from differentially expressed genes after stimulation of HUVEC with TNF- α compared with non-stimulated cells. Furthermore, most of the aforementioned genes have been identified as regulated by more than one metabolite, while a

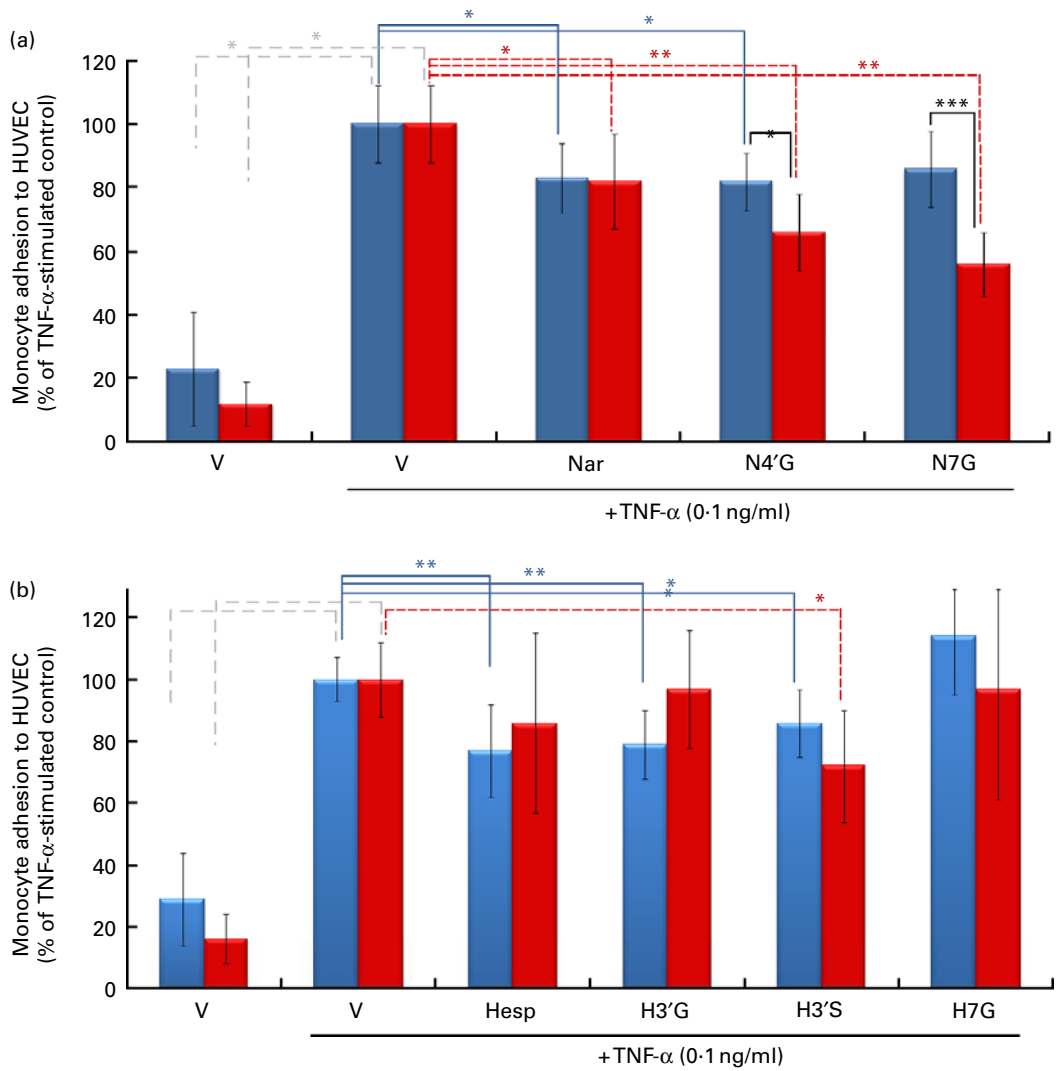


Fig. 3. Impact of pre-exposure of both endothelial cells and monocytes to flavanones on cell adhesion. Human umbilical vein endothelial cells (HUVEC) were stimulated with TNF- α for 4 h after incubation without or with (a) naringenin (Nar) and its conjugates (naringenin-4'-glucuronide (N4'G) and naringenin-7-glucuronide (N7G)) or (b) hesperetin (Hesp) and its conjugates (hesperetin-3'-glucuronide (H3'G); hesperetin-3'-sulphate (H3'S) and hesperetin-7-glucuronide (H7G)) at 2 μ M for 24 h. Prior to the adhesion assay, monocytes (U937) were pre-incubated or not with (a) Nar and its conjugates or with (b) Hesp and its conjugates at 2 μ M for 24 h, co-incubated for 1 h and their adhesion to HUVEC investigated. Monocyte adhesion is presented as a percentage of U937 to HUVEC. Values are means of three independent experiments, with standard deviations represented by vertical bars. Mean values were significantly different (by one-way ANOVA followed by Dunnett's *post hoc* test): * $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$. ■, Exposure to HUVEC only; ■, exposure to both HUVEC and monocytes. (A colour version of this figure can be found online at <http://www.journals.cambridge.org/bjn>)

few of them, such as *KEAP1* or *CCL2*, have been identified as modulated by only one of the studied metabolites.

The obtained expression profiles of the studied genes have been clustered using the hierarchical unsupervised method using PermutMatrix software. In this cluster, vertical lines represent different experimental conditions, while horizontal lines present the studied genes (Fig. 4). The cluster revealed two distinct groups, one corresponding to the expression profile of cells stimulated with TNF- α , noted A, and the second one containing the expression profiles of cells pre-exposed to flavanones and then stimulated with TNF- α , noted B. This cluster revealed that TNF- α stimulation of HUVEC induced up-regulation of most of the genes, while pre-exposure of HUVEC to flavanones results in a global down-regulation of gene expression.

Effect of flavanones and their metabolites on cell adhesion protein expression levels in TNF- α -stimulated endothelial cells

Together with gene expression analysis, protein expression was also analysed for flavanones and their metabolites that significantly reduced monocyte adhesion to HUVEC at 2 μ M concentration (H, H3'G, H3'S, naringenin and N4'G). Western blot analysis was performed for cell adhesion molecule proteins (ICAM1, VCAM1 and E-selectin) and cytoskeleton organisation protein (vinculin). The treatment of HUVEC with H, H3'S and naringenin (2 μ M) for 24 h prior to exposure to TNF- α induced a significantly decreased protein level of vinculin (Fig. 5), whereas no impact of the tested metabolites was observed on cell adhesion protein levels (Fig. S2, available online).

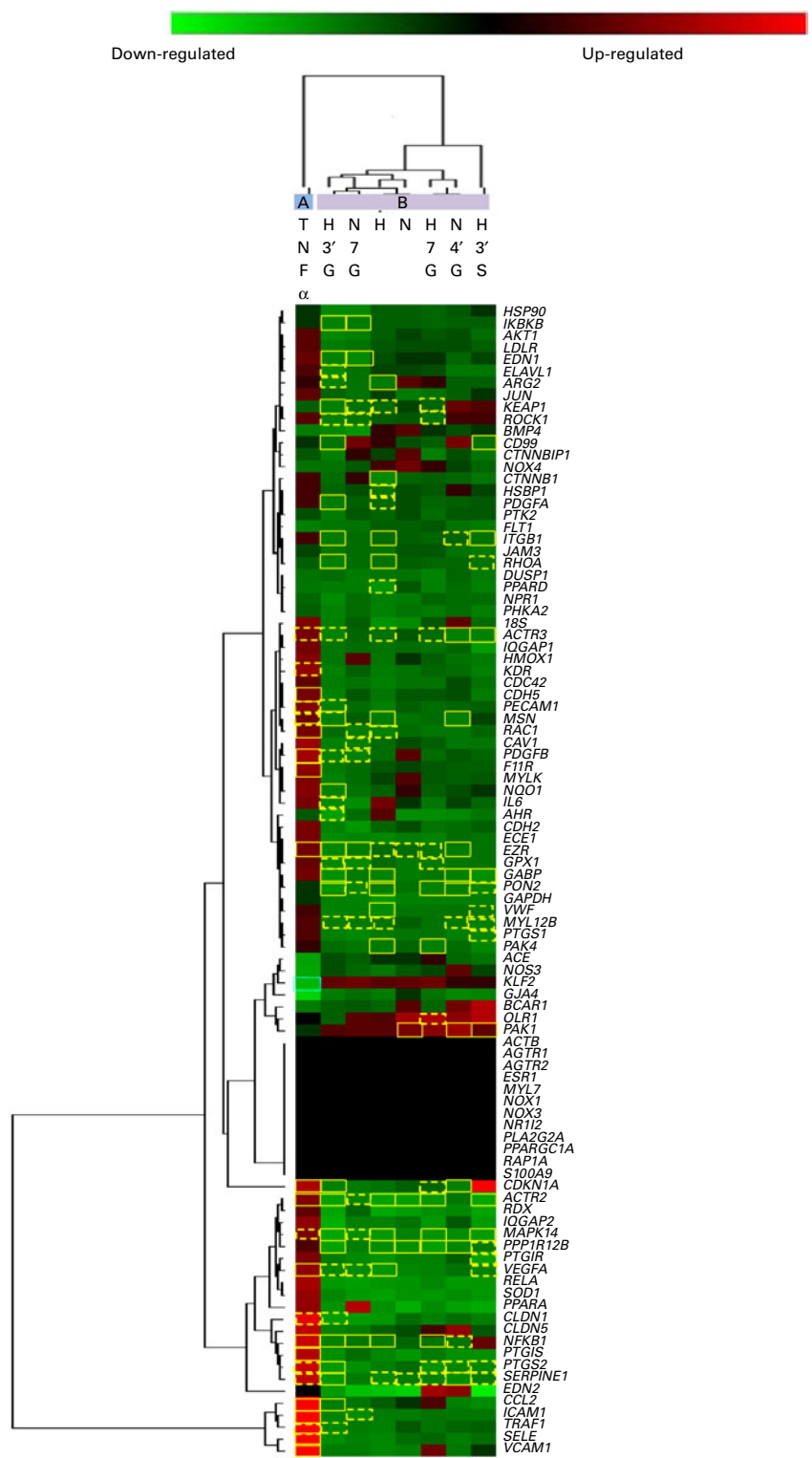


Fig. 4. Two-dimensional hierarchical clustering analysis of mRNA profiles in human umbilical vein endothelial cells (HUVEC) under different conditions (TNF- α *v.* control, TNF- α -stimulated HUVEC treated with flavanones *v.* TNF- α alone-stimulated HUVEC). Flavanones, aglycones (hesperetin (H) and naringenin (N)) and metabolites (hesperetin-3'-glucuronide (H3'G), hesperetin-3'-sulphate (H3'S), hesperetin-7-glucuronide (H7G), naringenin-4'-glucuronide (N4'G) and naringenin-7-glucuronide (N7G)) were used at 2 μ M concentration. mRNA are represented on the vertical axis, while different tested conditions are plotted on the horizontal axis. Red or green colours indicate up-regulation and down-regulation, respectively. Results are from three independent experiments. Significance was determined by the comparative C_T ($\Delta\Delta C_T$) method. Yellow squares (□): $P < 0.05$, blue dotted squares (□): $P < 0.1$. See Appendix 1 and Table S1 (available online) for a list of names and abbreviations. (A colour version of this figure can be found online at <http://www.journals.cambridge.org/bjn>)

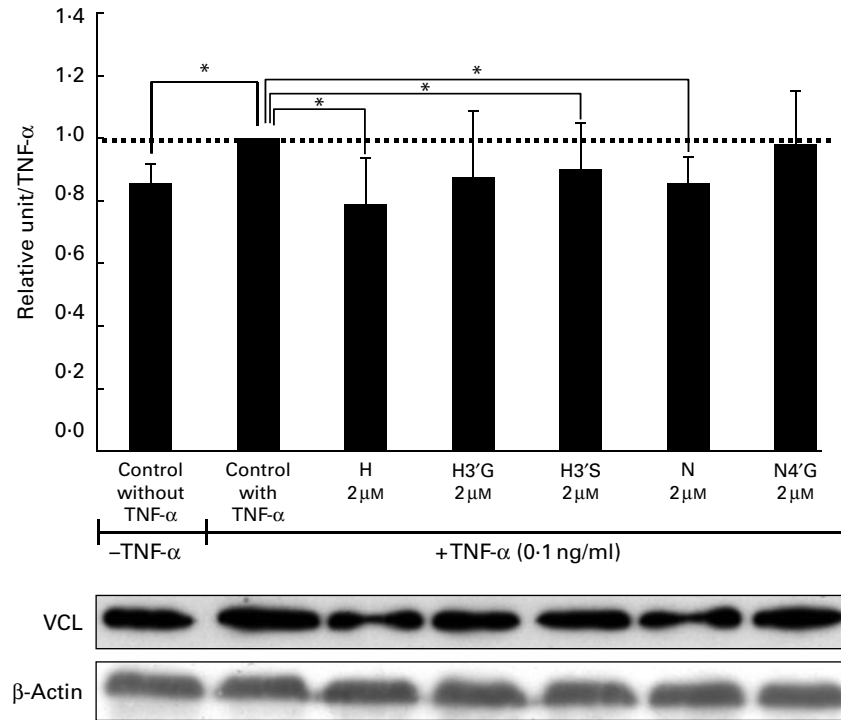


Fig. 5. Effect of flavanones and their metabolites on vinculin (VCL) protein expression in human umbilical vein endothelial cells (HUVEC). HUVEC were pre-treated with hesperetin (H), hesperetin-3'-glucuronide (H3'G), hesperetin-3'-sulphate (H3'S), naringenin (N) or naringenin-4'-glucuronide (N4'G) for 24 h followed by induction with TNF- α . Expression of VCL protein in cells was evaluated by immunoblotting. Values are means, with their standard errors represented by vertical bars, n 3. *Mean value was significantly different from that of TNF- α control without flavanone pre-treatment ($P < 0.05$).

Discussion

In a previous *in vivo* study, we demonstrated that flavanones consumed at a nutritional dose reduced lesion development in a mouse model of diet-induced atherosclerosis⁽²³⁾. Transcriptomic data analysis from aorta revealed that this effect might be mediated by flavanone-induced altered expression of genes that govern monocyte adhesion to endothelial cells and their subsequent trans-endothelial migration into the intima of aorta. Thus, to gain further insights into the role of flavanones on cardiovascular protection, the present study aimed to investigate the direct effects of circulating flavanone metabolites on endothelial cell function. Experimental conditions were set in order to reproduce the vascular environment observed after the consumption of flavanone-rich foods. From this perspective, flavanone metabolites that had been identified in human plasma after the consumption of orange and grapefruit juice were used. These metabolites were employed at physiologically relevant concentrations and were incubated with endothelial cells for 24 h, which corresponds to the time required for flavanones to be excreted following their ingestion⁽²³⁾. Furthermore, HUVEC were used as a model of human endothelial cells and were exposed to inflammatory stress to mimic a pro-atherogenic environment. To induce inflammatory stress, TNF- α , a pro-inflammatory cytokine capable of inducing endothelial dysfunction and whose circulating concentrations have been identified as elevated in plasma of patients at risk of CVD, was chosen⁽²⁴⁾. The present study showed that, except H7G and N7G, flavanone

metabolites were effective at 2 μ M to attenuate monocyte adhesion to TNF- α -activated endothelial cells, the first step of atherogenesis. This concentration has been reported in human plasma following grapefruit or orange consumption^(25,26) and can be therefore considered as nutritionally achievable. In the present experimental conditions, we failed to observe a dose-dependent effect of flavanone metabolites on cell adhesion. A similar observation has been previously reported for other flavonoids, both in endothelial and in immune cells^(27,28). Data from the present study suggest that metabolites could be as potent as their aglycones in preventing monocyte adhesion to activated HUVEC. Furthermore, it is noteworthy that flavanone metabolites conjugated on the B ring seem to be more efficient in reducing cell adhesion than those substituted on the A ring. This suggests that the biological potency of metabolites depends on the position of conjugation rather than on the nature of the substituent (sulphate or glucuronide). Overall, the ability of flavanone metabolites to decrease monocyte adhesion to activated endothelial cells may be one of the mechanisms underlying the anti-atherogenic activity of flavanones *in vivo*^(10–15).

When present in blood circulation, flavanone metabolites may interact with endothelial cells as well as with immune cells, such as monocytes. Consequently, to fit as closely as possible with physiological conditions, it has been examined to what extent the pre-exposure of both endothelial cells and monocytes to flavanone metabolites may affect the cell adhesion process. The obtained data showed that exposure of monocytes to N4'G and N7G potentiated the inhibitory

effect of flavanones on monocyte adhesion to HUVEC when compared with a condition in which only HUVEC were pre-exposed to metabolites (Fig. 3). This finding revealed that circulating monocytes also constitute cellular targets for naringenin metabolites, which affect the ability of monocytes to interact with endothelial cells. In agreement, we have previously shown in human subjects that hesperidin supplementation modulated gene expression in leucocytes, in particular genes involved in chemotaxis, adhesion or infiltration⁽²⁹⁾.

Gene expression analysis in endothelial cells using TaqMan low-density arrays suggested that flavanones could affect endothelial function through transcriptional regulation of gene expression. Genes whose expression was identified as modulated by TNF- α are involved in different cellular processes, such as cell adhesion (ICAM-1 and CDH5), inflammation and chemotaxis (NFKB1 and CCL2) or cytoskeleton organisation (ACTR2, EZR and RAC-1). This observed effect of TNF- α on gene expression has been previously reported⁽³⁰⁾. Exposure of endothelial cells to flavanones, aglycones or to their metabolites resulted in changes in expression of genes. Most of the modulated genes are involved in various cellular processes, including cell adhesion, cytoskeleton organisation, inflammation and chemotaxis, all being related to atherogenesis. Clustering of gene expression profiles revealed a specific pattern of gene expression in HUVEC activated by TNF- α . Interestingly, genes whose expression was up-regulated by TNF- α seemed to be down-regulated by flavanones and vice versa (Fig. 4). This observation revealed the potency of flavanone metabolites, at physiologically relevant concentrations, to counteract the effect induced by an inflammatory stimulus (TNF- α). As a result, genes related to inflammation could be the molecular targets associated with the reduction of monocyte adhesion to endothelial cells exposed to flavanone metabolites.

The inflammation process plays a critical role in atherosclerosis development through the recruitment of immune cells to the inflammatory site, particularly by triggering the synthesis of adhesion and chemotactic molecules. The present study showed that exposure of endothelial cells to flavanones, aglycones or metabolites induced an anti-inflammatory profile, as revealed by a down-regulation of expression of genes coding for inflammatory mediators, genes whose expression has been induced by TNF- α . *NFKB1* gene expression was down-regulated by H3'S, N7G, H and H7G compared with the TNF- α -stimulated control. NFKB1 encodes for a NF- κ B subunit, the well-known transcription factor involved in inflammation and activated by TNF- α ⁽³¹⁾. In agreement with the decreased expression of *NFKB1* gene in response to flavanones, a trend towards the down-regulation of some of its target genes, like *CCL2* (also called monocyte chemoattractant protein-1 and involved in the chemotaxis of monocytes to endothelial cells) or *Traf1* (one member of the TNF- α receptor)^(32,33) is observed. In addition, pre-exposure of endothelial cells to flavanones, aglycones or metabolites also affected the expression of genes coding for adhesion molecules, which constitute other targets of NF- κ B. For example, flavanones decreased expression of genes coding for adhesion

molecules like CD99 or ITGB1 and tended to reduce the expression of genes coding for E-selectin, ICAM-1 and VCAM-1 when compared with the TNF- α -stimulated control. These adhesion molecules participate in the rolling and firm adhesion of leucocytes to activated endothelium, and their down-regulation result in the impairment of monocyte adhesion to endothelial cells⁽¹⁴⁾. Overall, these results suggest that flavanones could inhibit inflammatory process through changes in the expression of inflammation-related genes. These genomic effects of flavanones could be, at least partially, mediated by their capacity to affect the expression of the NF- κ B transcription factor. This hypothesis is supported by other studies which have previously shown the ability of flavanones to suppress NF- κ B activation and related gene expression in rat kidneys or activated macrophages^(34–36).

Leucocyte adhesion on activated endothelium promotes the remodelling of the apical endothelial plasma membrane into actin-rich projections that surround adherent leucocytes, known as a docking structure⁽³⁷⁾. These trans migratory cups participate in firm adhesion of monocytes to the endothelial cells and serve as directional guides for leucocytes to cross the endothelium⁽³⁸⁾. These features of the actin cytoskeleton are also regulated by a cohort of actin-associated cytoskeletal proteins, such as talin, vinculin or cofilin. The complex constituted by ICAM-1 and VCAM-1 associated to ERM proteins (EZR, radixin and moesin) initiates intracellular signalling, such as activation of the small GTPase pathway^(37,39). Interestingly, TLDA analysis revealed that, when compared with the TNF- α -stimulated control, pre-exposure of endothelial cells to flavanones also down-regulated the expression of genes coding for ERM proteins: EZR, moesin and small GTPase members. In response to flavanones, the down-regulation of EZR reached significance for H3'S, N7G and N4'G and that of moesin for H3'S, H and N4'G. Pre-exposure of HUVEC to flavanones also resulted in a decreased expression of RAC1, a member of the rho family of small GTPase and whose inhibition reduces leucocyte extravasation^(37,39,40). RAC signalling is known to lead to actin polymerisation through the activation of actin-related protein (ARP) 2/3 complex⁽⁴¹⁾, which in turn allows transmigration⁽⁴²⁾. Interestingly, the present study also showed that most of the studied flavanone compounds significantly down-regulated the expression of both *ARP2* and *ARP3* genes coding for ACTR2 and ACTR3 proteins, which are major constituents of the ARP2/3 complex. Furthermore, gene expression analysis revealed differentially expressed genes involved in cytoskeleton organisation. Using Western blot analysis, we examined expression of vinculin, a protein involved in cytoskeleton organisation and remodelling. This analysis revealed a decrease in vinculin protein level in endothelial cells pre-exposed to flavanone metabolites (Fig. 5). Overall, the reduced gene expression of actin cytoskeleton, docking structure components, cytoskeleton organisation or their regulators by flavanones (aglycones and their metabolites) might result in decreased trans migratory cups formation and consequently lead to a lower monocyte adhesion to endothelial cells and transendothelial migration.

In conclusion, the present study reveals that at physiologically achievable concentrations, the flavanone metabolites present in human plasma following citrus fruit consumption retain the ability to decrease monocyte adhesion to TNF- α -stimulated endothelial cells. These metabolites also present potency to modulate expression of genes, particularly those coding for cell adhesion molecules, chemokines and cytoskeleton components that are involved in monocyte adhesion and recruitment in the vascular wall. The observed modulation of gene expression at the transcriptional level by flavanones might be related to the decreased adhesion of monocytes to endothelial cells. These data give an insight into potential cellular and molecular mechanisms responsible for the athero-protective effects of dietary flavanones reported *in vivo*.

Supplementary material

To view supplementary material for this article, please visit <http://dx.doi.org/10.1017/S0007114512005454>

Acknowledgements

We would like to thank Christiane Deval for TLDA experiments and Nicolas Gerard for Western blot analysis. We thank UNIJUS, the French association of fruit juice producers, who followed and communicated about this research programme. A. C. contributed to the conception and design of the study, acquisition, analysis and interpretation of data; writing the article and the final approval of the manuscript. D. M. contributed to the conception and design of the study, acquisition, analysis and interpretation of gene expression data; writing the article and the final approval of the manuscript. S. C. contributed to the Western blot analyses, writing the article and the final approval of the manuscript. J. A. M. M. contributed to the conception of the study, analysis and interpretation of cell adhesion data; writing the article and the final approval of the manuscript. M. K. K. contributed to the conception of the study, production and analysis of metabolites and the final approval of the manuscript. N. R. contribution to the conception of the study, production and analysis of metabolites and the final approval of the manuscript. S. S. contributed to the conception of the study, production and analysis of metabolites and the final approval of the manuscript. A. M. B. contributed to the development of the study protocol, revising of the article and the final approval of the manuscript. C. B.-P. contributed to the development of the study protocol, critical revising of the article and the final approval of the manuscript. A. M. contributed to the development of the study protocol, statistical analysis of the data and the final approval of the manuscript. C. M. contributed to the conception and design of the study, analysis and interpretation of data; writing the article and the final approval of the version to be published. The authors do not have financial/commercial conflict of interest. This project has received funding from the French National Research Agency in the context of the National Program for Research on Food and Nutrition (PNRA).

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Appendix 1. Gene abbreviations and names

<i>18S</i>	RNA, 18S ribosomal 0
<i>ACE</i>	Angiotensin I converting enzyme
<i>ACTB</i>	Actin, beta
<i>ACTR2</i>	Actin-related protein 2
<i>ACTR3</i>	Actin-related protein 3
<i>AGTR1</i>	Angiotensin II receptor, type 1
<i>AGTR2</i>	Angiotensin II receptor, type 2
<i>AHR</i>	Aryl hydrocarbon receptor
<i>AKT1</i>	v-akt murine thymoma viral oncogene homolog 1
<i>ARG2</i>	Arginase, type II
<i>BCAR1</i>	Breast cancer anti-estrogen resistance protein 1
<i>BMP4</i>	Bone morphogenetic protein 4
<i>CAV1</i>	Caveolin 1
<i>CCL2</i>	Hemokine (C-C motif) ligand 2
<i>CD99</i>	CD99 molecule
<i>CDC42</i>	Cell division control protein 42 homolog
<i>CDH2</i>	Cadherin-2
<i>CDH5</i>	Cadherin-5
<i>CDKN1A</i>	Cyclin-dependent kinase inhibitor 1
<i>CLDN1</i>	Claudin 1
<i>CLDN5</i>	Claudin 5
<i>CTNNB1</i>	Catenin (cadherin-associated protein), beta 1, 88 kDa
<i>CTNNBIP1</i>	Catenin, beta interacting protein 1
<i>DUSP1</i>	Dual specificity phosphatase 1
<i>ECE1</i>	Endothelin converting enzyme 1
<i>EDN1</i>	Endothelin 1
<i>EDN2</i>	Endothelin 2
<i>ELAVA1</i>	ELAV (embryonic lethal, abnormal vision, Drosophila)-like 1 (Hu antigen R)
<i>ESR1</i>	Estrogen receptor 1
<i>EZR</i>	Ezrin
<i>F11R</i>	F11 receptor
<i>FLT1</i>	Fms-related tyrosine kinase 1
<i>GABP</i>	GA binding protein transcription factor
<i>GAPDH</i>	Glyceraldehyde-3-phosphate dehydrogenase
<i>GJA4</i>	Gap junction protein, alpha 4, 37 kDa
<i>GPX1</i>	Glutathione peroxidase 1
<i>HMOX1</i>	Heme oxygenase (decycling) 1
<i>HSBP1</i>	Heat shock factor binding protein 1
<i>HSP90</i>	Heat shock protein 90
<i>ICAM1</i>	Intercellular adhesion molecule 1
<i>IKKBK</i>	Inhibitor of kappa light polypeptide gene enhancer in B-cells, kinase beta
<i>IL6</i>	Interleukin 6
<i>IQGAP1</i>	IQ motif containing GTPase activating protein 1
<i>IQGAP2</i>	IQ motif containing GTPase activating protein 2
<i>ITGB1</i>	Integrin, beta 1
<i>JAM3</i>	Junctional adhesion molecule 3
<i>JUN</i>	Jun proto-oncogene
<i>KDR</i>	Kinase insert domain receptor
<i>KEAP1</i>	Kelch-like ECH-associated protein 1
<i>KLF2</i>	Kruppel-like factor 2
<i>LDLR</i>	Low density lipoprotein receptor
<i>MAPK14</i>	Mitogen-activated protein kinase 14
<i>MSN</i>	Moesin
<i>MYL12B</i>	Myosin, light chain 12B
<i>MYL7</i>	Myosin, light chain 7
<i>MYLK</i>	Myosin light chain kinase
<i>NFKB1</i>	Nuclear factor of kappa light polypeptide gene enhancer in B-cells 1
<i>NOS3</i>	Nitric oxide synthase 3
<i>NOX1</i>	NADPH oxidase 1
<i>NOX3</i>	NADPH oxidase 3
<i>NOX4</i>	NADPH oxidase 4
<i>NPR1</i>	Natriuretic peptide receptor A/guanylate cyclase A

Appendix 1. Continued

<i>NQO1</i>	NAD(P)H dehydrogenase, quinone 1
<i>NR1I2</i>	Nuclear receptor subfamily 1, group 1, member 1
<i>OLR1</i>	Oxidized low density lipoprotein (lectin-like) receptor 1
<i>PAK1</i>	p21 protein (Cdc42/Rac)-activated kinase 1
<i>PAK4</i>	p21 protein (Cdc42/Rac)-activated kinase 4
<i>PDGFA</i>	Platelet-derived growth factor alpha polypeptide
<i>PDGFB</i>	Platelet-derived growth factor beta polypeptide
<i>PECAM1</i>	Platelet/endothelial cell adhesion molecule 1
<i>PHKA2</i>	Phosphorylase kinase, alpha 2
<i>PLA2G2A</i>	Phospholipase A2, group IIA
<i>PON2</i>	Paraoxonase 2
<i>PPARA</i>	Peroxisome proliferator-activated receptor alpha
<i>PPARD</i>	Peroxisome proliferator-activated receptor delta
<i>PPARGC1A</i>	Peroxisome proliferator-activated receptor gamma, coactivator 1 alpha
<i>PPP1R12B</i>	Protein phosphatase 1, regulatory subunit 12B
<i>PTGIR</i>	Prostaglandin I2 (prostacyclin) receptor
<i>PTGIS</i>	Prostaglandin I2 (prostacyclin) synthase
<i>PTGS1</i>	Prostaglandin-endoperoxide synthase 1
<i>PTGS2</i>	Prostaglandin-endoperoxide synthase 2
<i>PTK2</i>	Protein tyrosine kinase 2
<i>RAC1</i>	Ras-related C3 botulinum toxin substrate 1
<i>RAP1A</i>	Ras-related protein Rap-1A
<i>RDX</i>	Radixin
<i>RELA</i>	v-rel reticuloendotheliosis viral oncogene homolog A
<i>RHOA</i>	Ras homolog family member A
<i>ROCK1</i>	Rho-associated, coiled-coil containing protein kinase 1
<i>S100A3</i>	S100 calcium binding protein A3
<i>SELE</i>	Selectin E
<i>SERPINE1</i>	Serpin peptidase inhibitor, clade E 1
<i>SOD1</i>	Superoxide dismutase 1
<i>TRAF1</i>	TNF receptor-associated factor 1
<i>VCAM1</i>	Vascular cell adhesion molecule 1
<i>VEGFA</i>	Vascular endothelial growth factor A
<i>VWF</i>	von Willebrand factor