Effect of quercetin and its metabolite on caveolin-1 expression induced by oxidized LDL and lysophosphatidylcholine in endothelial cells

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Oxidized low-density lipoprotein contributes to atherosclerotic plaque formation, and quercetin is expected to exert anti-atherosclerotic effects. We previously reported accumulation of conjugated quercetin metabolites in the aorta of rabbits fed high-cholesterol diets with quercetin glucosides, resulting in attenuation of lipid peroxidation and inhibition of lipid accumulation. Caveolin-1, a major structural protein of caveolae in vascular endothelial cells, plays a role in atherosclerosis development. Here we investigated effects of oxidized low-density lipoprotein, quercetin and its metabolite, quercetin 3-O-β-glucuronide, on caveolin-1 expression. Oxidized low-density lipoprotein significantly upregulated caveolin-1 mRNA expression. An oxidized low-density lipoprotein component, lysophosphatidylcholine, also induced expression of both caveolin-1 mRNA and protein. However, lysophosphatidylcholine did not affect the location of caveolin-1 proteins within caveolae structures. Co-treatment with quercetin or quercetin 3-O-β-glucuronide inhibited lysophosphatidylcholine-induced caveolin-1 expression. Quercetin and quercetin 3-O-β-glucuronide also suppressed expression of adhesion molecules induced by oxidized low-density lipoprotein and lysophosphatidylcholine. These results strongly suggest lysophosphatidylcholine derived from oxidized low-density lipoprotein contributes to atherosclerotic events by upregulating caveolin-1 expression, resulting in induction of adhesion molecules. Quercetin metabolites are likely to exert an anti-atherosclerotic effect by attenuating caveolin-1 expression in endothelial cells.

Key Words: caveolin-1, adhesion molecule, quercetin, quercetin metabolite, vascular endothelial cell

Quercetin (3,3′,4′,5,7-pentahydroxyflavone; Q), a typical flavonol-type flavonoid, frequently presents as its glycoside form in fruits and vegetables. Epidemiological studies indicate dietary intake of flavonoids, including Q, is inversely associated with risk of cardiovascular diseases such as coronary heart disease(1,2) and stroke.(3) A Japanese study reported that intake of flavonoids, in particular Q, inversely correlated with plasma total cholesterol and low-density lipoprotein (LDL) concentrations.(4) In addition to reducing oxidative stress, dietary Q appears to exert effects on vascular function by elevating both endothelial nitric oxide synthase expression and blood glutathione redox ratio.(5)

Dietary Q is converted to glucuronidate and/or sulfate derivatives, or their Oxo-methyl derivatives, during absorption into the body, and presents exclusively as these conjugated metabolites in circulating blood.(6) In a previous study utilizing high-cholesterol fed rabbits, we reported that dietary intake of Q glucoside resulted in accumulation of its conjugated metabolites within the aorta, leading to inhibition of aortic lipid peroxidation and accumula-tion.(7) We also demonstrated that Q 3-O-β-glucuronide (Q3GA), one of Q’s major conjugated metabolites, can return to its aglycone form to exert physiological functions in activated macrophages, an event expected to be involved in early atherosclerosis development.(8,9) Although oxidative damage to endothelial cells may profoundly affect vascular functions, effects of Q metabolites on vascular endothelial cells remain obscure.

Circulating LDL is oxidized in intima and resulting oxidized LDL (ox-LDL) plays diverse roles in atherosclerosis pathogenesis and progression.(10) In the early stages of atherosclerotic plaque formation, ox-LDL activates endothelial cells by inducing cell surface adhesion molecules, especially intracellular cell adhesion molecule-1 (ICAM-1) and vascular cell adhesion molecule-1 (VCAM-1), which mediate rolling and adhesion of monocytes.(11) Caveolae, a lipid raft subtype, are 50–100 nm flask-shaped invaginations of plasma membrane, which are abundant in endothelial cells. Caveolin-1 (CAV-1) is a major structural protein of caveolae. (12) Endothelial-specific over-expression of CAV-1 accelerates progression of atherosclerosis by reducing endothelial cell proliferation, migration and nitric oxide (NO) production leading to VCAM-1 expression.(13) Caveolae and CAV-1 mediate LDL uptake/transcytosis and regulate LDL entry into arterial walls.(14,15) Furthermore, it has been suggested that CAV-1 directly regulates expression of these adhesion molecules.(15) These facts suggest CAV-1 plays a critical role in regulation of endothelial cell activation. Therefore, we aimed to clarify effects of Q metabolites on ox-LDL-induced CAV-1 expression and adhesion molecule expression in human umbilical vein endothelial cells (HUVEC).

Materials and Methods

Reagents. Quercetin dehydrate, α-tocopherol, L-α-lysophosphatidylcholine (lysoPC) and rabbit polyclonal antibody to CAV-1 were purchased from Sigma-Aldrich (St. Louis, MO). Human LDL and human oxidized-LDL (low-thiobarbituric acid reactive) were obtained from Biomedical Technologies (Stoughton, MA). Q3GA was obtained from Extrasynthese (Genay, France). 13-Hydroperoxyoctadecadienoic acid (13-HPODE) was obtained from Larodan Fine Chemicals AB (Malmo, Sweden). 13(S)-Hydroxyoctadecadienoic acid (13-HODE) was obtained from Cayman Chemical (Ann Arbor, MI). 4-Hydroxynonenal (4-HNE) was obtained from Percipio Biosciences (Burlingame, CA). Rabbit polyclonal antibody to glyceraldehyde-3-phosphate dehydrogenase (GAPDH) and anti-phospho-CAV-1 antibody were obtained from 

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Cell Signaling Technology (Beverly, MA). Secondary antibody, polyclonal goat anti-rabbit immunoglobulin/horseradish peroxidase was obtained from Dako (Glostrup, Denmark).

**Cells and culture.** HUVEC (Lonza, Basel, Switzerland), were cultured in endothelial cell growth medium, EGM-2 (Lonza), and maintained at 37°C in a humidified atmosphere containing 5% CO₂. Cells were used in experiments from passages 3–10.

**Real-time reverse-transcription polymerase chain reaction (RT-PCR).** HUVEC were plated into 24-well plates (4 × 10⁵ cells/well) and incubated for 24 h. Cells were treated with ox-LDL or related compounds, as described in figure legends. To estimate effects of flavonoids, they were added to cells at the same time as lipids or related compounds. In 6 h pretreatment experiments, flavonoids were washed out with Hank’s balanced salt solution buffer, which was replaced with fresh medium containing lipids or related compounds. Total RNA was isolated from HUVEC using ISOGEN (Nippon Gene, Toyama, Japan). mRNA expression was determined by real-time RT-PCR, as previously described. Primers used to examine genes are listed in Table 1. Primers for CAV-1 and GAPDH were obtained from Takara Bio, Inc. (Otsu, Japan). Primers for ICAM-1 and VCAM-1 were synthesized according to previous report.

Reaction conditions for RT and PCR were based on protocols provided by Applied Biosystems (Foster City, CA). Relative levels of gene expression for each sample were calculated using a comparative Ct method. Expression of target genes (CAV-1, ICAM-1 and VCAM-1) in each sample was normalized to GAPDH Ct values. Data are expressed as mean ± SD of three separate experiments.

**Cell viability assay.** HUVEC were plated into 96-well plates (2 × 10⁴ cells/well). After a 24 h incubation, culture medium was replaced with 100 μl of fresh medium. Ox-LDL-related lipids were dissolved in ethanol and added to culture medium, followed by incubation for an additional 24 h. Cells were treated with 5 μl of Cell Proliferation Reagent WST-1 (Roche, Cat No. 11644807001, Shire, UK). Images were captured with a Lumino Image Analyzer LAS-3000 mini (Fujiﬁlm, Tokyo, Japan).

**Table 1. Primers used for real-time RT-PCR**

<table>
<thead>
<tr>
<th>Gene</th>
<th>Forward</th>
<th>Reverse</th>
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<tbody>
<tr>
<td>CAV-1</td>
<td>5'-TCTGGCTTACCTCAGGCAAC-3'</td>
<td>5'-GCTAGCCATTATGTGCC-3'</td>
</tr>
<tr>
<td>ICAM-1</td>
<td>5'-TCTGGTCCCCTCAAAAGTCT-3'</td>
<td>5'-GGGTCCTATGGCACAACA-3'</td>
</tr>
<tr>
<td>VCAM-1</td>
<td>5'-GCTGCTGATGAGCTGACAT-3'</td>
<td>5'-GCTCAGAGGCTGCTATAC-3'</td>
</tr>
<tr>
<td>GAPDH</td>
<td>5'-GCACCAGCAAGGCTGGAAC-3'</td>
<td>5'-TGGTGAAGACGCCGATGCA-3'</td>
</tr>
</tbody>
</table>

**Partion of microdomains and non-microdomains in cell membranes.** HUVEC were seeded into 60-mm dishes (4 × 10⁴ cells/dish). After incubation for 24 h, cells were treated with lysophPC or methyl-beta-cyclodextrin (MβCD). Cell lysates were subjected to ultracentrifugation and fractionated as previously described. Isolation of microdomains was confirmed by dot blots, and CAV-1 of each fraction was detected.

**Statistical analyses.** Data are expressed as mean ± SD from at least three independent experiments. Statistical analyses were performed using PASW Statistics 18.0 (IBM, Armonk, NY). Data were analyzed by one-way analysis of variance with Tukey multiple comparison test (p<0.05).

**Results**

**Effect of ox-LDL on CAV-1 mRNA expression and suppression by Q and Q3GA.** CAV-1 mRNA expression was upregulated by addition of ox-LDL in the range of 10–100 μg/ml, in a concentration-dependent manner (Fig. 1). In contrast, LDL itself did not exert such an effect. Next, HUVEC were exposed to ox-LDL (100 μg/ml) simultaneously with Q (1, 10 μM) or Q3GA (1, 10 μM) and incubated for 12 h to examine effects of Q and Q3GA on ox-LDL-induced CAV-1 mRNA expression. Significant suppression of ox-LDL-induced CAV-1 mRNA expression was observed with co-administration of 10 μM Q (Fig. 1). Q3GA also tended to suppress CAV-1 mRNA expression, although this effect was not significant.

**Effect of lysoPC and ox-LDL-related lipid peroxidation products on CAV-1 mRNA expression.** First, toxicity of lysoPC and lipid peroxidation products derived from ox-LDL was examined using a WST-1 assay (Fig. 2A). While administration of 0.1 or 1 μM lysoPC did not affect cell viability, 1 μM 4-HNE decreased cell viability by 55%. Neither 13-HPODE nor 13-HODE affected cell viability, even at 10 μM. For our treatment condition, we decided to use a concentration showing non-cytotoxicity. Therefore, concentrations of each compound varied in the following experiments. CAV-1 mRNA expression was upregulated by lysoPC at 0.1 μM, but not 1 μM, whereas 0.1 μM 4-HNE had no effect on expression. 13-HPODE, but not 13-HODE, upregulated CAV-1 mRNA expression at 10 μM (Fig. 2B). CAV-1 mRNA expression induced by 0.1 μM lysoPC increased in a time-dependent manner for 24 h (Fig. 2C); however, lysoPC did not induce phosphorylation of CAV-1 (Fig. 2D).

**Effect of lysoPC on CAV-1 distribution in cell membranes.** Effect of lysoPC on CAV-1 distribution in cell membrane lipid rafts was examined using an ultracentrifugation technique (Fig. 2E). In control cells (only treated with vehicle), CAV-1 existed in the third and fifth fractions (lipid raft region). MβCD, which disrupts lipid rafts by removing cholesterol from cell membranes, broadened observance of CAV-1 through the
seventh fraction. After incubation of HUVEC with 0.1 μM lysoPC, distribution of CAV-1 remained unchanged in the lipid raft region (third to fifth fractions). Thus, it was confirmed that lysoPC did not affect location of CAV-1 in cellular membranes.

Effect of Q and Q3GA on lysoPC-induced CAV-1 expression. Both Q and Q3GA suppressed lysoPC-induced CAV-1 mRNA expression (Fig. 3A). CAV-1 protein levels were upregulated by addition of lysoPC for 48 h (Fig. 3B), whereas 10 μM Q suppressed upregulation of CAV-1 protein levels. Q3GA also tended to suppress CAV-1 protein upregulation, although this was not significant. To avoid direct interaction between Q/Q3GA and lysoPC in culture medium, Q or Q3GA were pre-incubated with cells for 6 h before addition of lysoPC. After removal of flavonoids from culture medium, cells were treated with lysoPC. Pre-incubation experiments also indicated that Q and Q3GA suppressed lysoPC-induced CAV-1 mRNA expression (Fig. 3C).

Effect of α-tocopherol on lysoPC-induced CAV-1 mRNA expression. We used α-tocopherol as a lipophilic antioxidant capable of exerting function in biomembranes. Fig. 4 shows co-administration of both 1 and 10 μM α-tocopherol suppressed lysoPC-induced CAV-1 mRNA expression.

Effect of Q and Q3GA on lysoPC- or ox-LDL-induced ICAM-1 and VCAM-1 mRNA expression. We first confirmed that mRNA expression of both ICAM-1 and VCAM-1 were upregulated by ox-LDL in a time-dependent manner with 24 h treatment (Fig. 5). After 18 h of incubation with lysoPC, expression of both adhesion molecules was significantly upregulated. Treatment with lysoPC for 24 h significantly upregulated ICAM-1 mRNA expression (Fig. 6A), which was significantly suppressed by Q and Q3GA in a concentration-dependent manner in the range examined (between 0.1 and 10 μM; Fig. 6A). VCAM-1 mRNA expression tended to be upregulated by lysoPC, although no significant difference was observed compared with controls at 24 h (Fig. 6A).

Both ICAM-1 and VCAM-1 mRNA expression were upregulated by addition of ox-LDL, but LDL itself did not show an elevating effect (Fig. 6B). Both Q and Q3GA were found to significantly suppress expression of ICAM-1 and VCAM-1 induced by ox-LDL (Fig. 6B).

Discussion

Results from our previous study using cholesterol-fed rabbits suggested dietary Q accumulates as conjugated metabolites within the aorta, where it attenuates lipid peroxidation and hyperlipidemia. Our immunohistochemical study using a monoclonal antibody demonstrated that Q metabolites target activated macrophages, resulting in selective deposition in human atherosclerotic arteries. Nevertheless, human epidemiological and interventional studies imply dietary Q exerts multiple mechanisms of action at diverse targets to protect vascular tissues from atherosclerosis. In this study, we focused on vascular endothelial cells as an alternative target of dietary Q. Endothelial activation and dysfunction are an initial event in atherosclerotic plaque formation, and endothelial CAV-1 plays a proatherogenic role. Furthermore, ox-LDL plays a critical role in pathogenesis and progression of atherosclerosis by several mechanisms including upregulation of CAV-1 expression. However, little is known about the effects of dietary Q on ox-LDL-induced endothelial activation through CAV-1 expression. Therefore, we investigated the effect of ox-LDL on CAV-1 expression, as well as the modulating effect of Q and its major metabolite, Q3GA, on ox-LDL-induced CAV-1 expression in HUVEC.

CAV-1-knockout mice showed lower expression of adhesion molecules, VCAM-1 and ICAM-1, observed transgenic mice overexpressing CAV-1 in endothelial cells have increased expression of VCAM-1. Fernandez-Hernando et al. observed CAV-1 co-localizes with ICAM-1 in caveolae structures and modulates ICAM-1-dependent translocation of monocytes. In addition, an Src family kinase responsible for CAV-1 signaling activates downstream elements of the ICAM-1 pathway, resulting in ICAM-1 expression throughout caveolae structure.
Fig. 2. Effect of lysoPC and ox-LDL-related lipid peroxidation products on CAV-1 mRNA expression and effect of lysoPC on CAV-1 distribution in cell membranes. (A) Cell viability in lysoPC or ox-LDL-related lipid peroxidation products-treated HUVECs. HUVECs were incubated with lysoPC, 4-HNE, 13-HODE or 13-HPODE for 24 h. (B) CAV-1 mRNA expression in HUVECs. HUVECs were incubated with ox-LDL related lipid for 24 h. (C) Effect of lysoPC (0.1 μM) on CAV-1 expression at 12, 18, and 24 h. (A–C) Values represent mean ± SD (n = 3). Letters represent significant differences among treatment groups. (D) Effect of 24 h lysoPC treatment on CAV-1 phosphorylation in HUVECs in the absence (−) or presence (+) of sodium orthovanadate (Na$_3$VO$_4$). (E) Effect of lysoPC on CAV-1 distribution in microdomain fractions in HUVECs. After incubation of HUVECs with lysoPC for 24 h or MβCD for 30 min, cell lysates were split into nine fractions and CAV-1 distribution was determined by dot blot analysis.
Fig. 3. Effect of Q and Q3GA on lysoPC-induced CAV-1 expression in HUVECs. (A) CAV-1 mRNA expression is induced in HUVECs by lysoPC and co-addition of Q/Q3GA for 24 h. (B) Western blot of CAV-1 and density of each image analyzed. HUVECs were incubated with lysoPC with Q or Q3GA for 48 h. (C) CAV-1 mRNA expression induced in HUVECs by 6 h preincubation with Q/Q3GA and subsequent incubation with lysoPC for 24 h. (A–C) Values represent mean ± SD (n = 3). Letters represent significant differences among treatment groups.
Hydroperoxides are lipid peroxidation products of ox-LDL that decompose into aldehydes such as 4-HNE. LysoPC is generated by an LDL-associated, platelet-activating factor-acylhydrolase-dependent hydrolysis of oxidized phosphatidylcholine. We previously reported that nonesterified fatty acid hydroperoxides released from phosphatidylcholine hydroperoxides can be reduced to hydroxy derivatives by the action of apoA1 present in HDL. Therefore, we selected lysoPC, 4-HNE, 13-HPODE and 13-HODE as representative ox-LDL-specific components and investigated their effects on CAV-1 expression in endothelial cells (Fig. 2A). Among them, 0.1 μM lysoPC and 10 μM 13-HPODE were found to induce CAV-1 expression without cell cytotoxicity. LysoPC is recognized to contribute to various atherosclerotic events. In endothelial cells, lysoPC is known to participate in expression of adhesion molecules ICAM-1, VCAM-1 and P-selectin. LysoPC appears to be a plausible endothelial cell activator, as lysoPC could induce CAV-1 expression at a lower concentration (0.1 μM) compared with 13-HPODE (10 μM). Thus, we deduced that lysoPC is at least partly responsible for the inductive effect of ox-LDL on CAV-1 expression.

Phosphorylation of CAV-1 is suggested to activate downstream signaling cascades associated with several endothelial functions. However, lysoPC exposure did not induce phosphorylation of CAV-1 in endothelial cells (Fig. 2D). Further, lysoPC did not affect localization of CAV-1 in caveolae structures, as distribution of CAV-1 was hardly changed by addition of lysoPC to cells (Fig. 2E). LysoPC reportedly destabilizes cellular membranes and/or acts as a disruptor. Thus, it was hypothesized that lysoPC affects function of CAV-1 by changing its location in caveolae structure. However, this hypothesis was ruled out and it can be concluded that lysoPC induces expression of CAV-1 without affecting caveolae.

LysoPC-induced gene expression of CAV-1 was effectively suppressed by simultaneous addition of Q or Q3GA (Fig. 3A). In vascular endothelial cells, lysoPC has been reported to promote O2- production via enhanced NADPH oxidase activity and induce mitochondrial reactive oxygen species (ROS) production. Here, a lipophilic antioxidant, α-tocopherol, also suppressed CAV-1 expression (Fig. 4), indicating antioxidant activity relates to attenuation of CAV-1 expression and subsequent adhesion molecule expression. CAV-1 expression in fibroblasts was promoted by hydrogen peroxide through activation of the p38 MAPK pathway, resulting from binding of specificity protein 1 (Sp1) to the promoter region, which increased transcription of CAV-1. Interestingly, Q has been shown to suppress promoter region activation and may suppress ROS-dependent signal transduction pathways by scavenging ROS generated during the incubation period. Pre-incubation of cells with Q or Q3GA before addition of lysoPC also resulted in effective inhibition of CAV-1 expression (Fig. 3C), suggesting a direct reaction of Q with lysoPC is not required to exert protective functions. It is noteworthy that both Q3GA and Q suppressed CAV-1 expression (Fig. 3A and C), although cellular uptake of Q3GA seems to barely occur because of its high hydrophilicity. Q3GA may localize at the surface of cell membranes to affect signal transduction pathway at the initial site.

LysoPC-stimulated ICAM-1 and VCAM-1 expression in a time-dependent manner (Fig. 5), although ICAM-1 increased more significantly than VCAM-1 after a 24 h incubation (Fig. 6A). LysoPC is known to promote expression of ICAM-1 and VCAM-1 at physiological concentrations, resulting in rolling and adhesion of monocytes. Activation of the PKC pathway may be involved in lysoPC-induced ICAM-1 expression in porcine coronary arterial endothelium. Enhancement of ICAM-1 expression was effectively suppressed by Q and Q3GA (Fig. 6A). Q appears to be more effective than Q3GA with regard to lysoPC-induced adhesion molecule expression. Although Q is scarcely present in circulating blood in its aglycone form, circulating conjugated Q metabolites may be deconjugated in the inflammatory site by

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**Fig. 4.** Effect of α-tocopherol on lysoPC-induced CAV-1 expression in HUVECs. HUVECs were incubated with lysoPC with α-tocopherol for 24 h. Values represent mean ± SD (n = 3). Letters represent significant differences among treatment groups.

**Fig. 5.** Effect of lysoPC treatment on ICAM-1 and VCAM-1 mRNA expression in HUVECs. HUVECs were incubated with lysoPC for indicated time (6, 12, 18 or 24 h). Values represent mean ± SD (n = 3). Letters represent significant differences among treatment groups.
efflux of β-glucuronidase from activated macrophages.\(^{(8,48)}\)

Resulting Q aglycone may act as a more effective suppressor of CAV-1 expression than Q metabolites upon exposure to lysoPC and other components present in ox-LDL. Koga and Meydani\(^{(49)}\) also suggested the aglycone form is required for Q to exert effective suppression of monocyte cell adhesion to human aortic endothelial cells. Green tea polyphenols and one of their components, epigallocatechin-3-gallate, were reported to protect endothelial cells from CAV-1 expression by modulating MAPK signaling.\(^{(50,51)}\) While it is expected that Q protects endothelial cells in a manner similar to green tea polyphenols, it should be noted that concentrations and structures required for effect vary greatly depending on each polyphenol. This study indicates effective concentrations of Q and Q3GA to modulate CAV-1 expression are around 1 μM, which seems to be within the physiological concentration of circulating blood.\(^{(52-54)}\) Thus, dietary Q may act as a modulator of vascular endothelial functions leading to antiatherosclerotic effects. This study may support results of previous interventional studies showing beneficial effects of flavonoid intake on reduction of cardiovascular disease risk and improvement of vascular endothelial function.\(^{(1-3,55)}\)

In conclusion, ox-LDL increased expression of CAV-1 mRNA, as well as expression of mRNA for adhesion molecules ICAM-1 and VCAM-1. Among ox-LDL-related components, lysoPC significantly stimulated CAV-1 and ICAM-1 expression, indicating lysoPC is partly responsible for the role of ox-LDL in mediating CAV-1 expression. Both Q and Q3GA suppressed ox-LDL and lysoPC-induced CAV-1 and adhesion molecule expression. Therefore, Q metabolites may exert anti-atherosclerotic effects by targeting CAV-1 expression in endothelial cells and dietary Q is a promising food factor to prevent atherosclerosis.

**Acknowledgments**

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Abbreviations

CAY-1 caveolin-1
GAPDH glyceraldehyde-3-phosphate
4-HNE 4-hydroxynonenal
13-HODE 13(S)-hydroxyoctadecadienoic acid
13-HPODE 13-hydroperoxyoctadecadienoic acid
HUVEC human umbilical vein endothelial cells
ICAM-1 intracellular cell adhesion molecule-1
LDL low-density lipoprotein
LysoPC lysophosphatidylcholine
MβCD methyl-beta-cyclodextrin
NO nitric oxide
ox-LDL oxidized low-density lipoprotein
Q quercetin
Q3GA quercetin-3-O-β-glucuronide
ROS reactive oxygen species
RT-PCR reverse-transcription polymerase chain reaction
VCAM-1 vascular cell adhesion molecule-1

Conflict of Interest

No potential conflict of interest were disclosed.

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