Intestinal Scavenger Receptors Are Involved in Vitamin K₁ Absorption*

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Background: Vitamin K intestinal absorption is poorly understood.

Results: Modulating SR-BI and CD36 expression or function in cells and in mice led to control of vitamin K₁ intestinal transport.

Conclusion: Intestinal scavenger receptors impact vitamin K₁ transport through the enterocyte.

Significance: Understanding vitamin K₁ intestinal absorption is crucial to maintain an optimal vitamin K status.

Vitamin K₁ (phylloquinone) intestinal absorption is thought to be mediated by a carrier protein that still remains to be identified. Apical transport of vitamin K₁ was examined using Caco-2 TC-7 cell monolayers as a model of human intestinal epithelium and in transfected HEK cells. Phylloquinone uptake was then measured ex vivo using mouse intestinal explants. Finally, vitamin K₁ absorption was compared between wild-type mice and mice expressing scavenger receptor class B type I (SR-BI) in the intestine and mice deficient in cluster determinant 36 (CD36). Phylloquinone uptake by Caco-2 cells was saturable and was significantly impaired by co-incubation with α-tocopherol (and vice versa). Anti-human SR-BI antibodies and BLT1 (a chemical inhibitor of lipid transport via SR-BI) blocked up to 85% of vitamin K₁ uptake. BLT1 also decreased phylloquinone apical efflux by ~80%. Transfection of HEK cells with SR-BI and CD36 significantly enhanced vitamin K₁ uptake, which was subsequently decreased by the addition of BLT1 or sulfo-N-succinimidyl oleate (CD36 inhibitor), respectively. Similar results were obtained in mouse intestinal explants. In vivo, the phylloquinone postprandial response was significantly higher, and the proximal intestine mucosa phylloquinone content 4 h after gavage was increased in mice expressing SR-BI compared with controls. Phylloquinone postprandial response was also significantly increased in CD36-deficient mice compared with wild-type mice, but their vitamin K₁ intestinal content remained unchanged. Overall, the present data demonstrate for the first time that intestinal scavenger receptors participate in the absorption of dietary phylloquinone.

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the mechanisms of this phenomenon have not been fully elucidated (12). We suggest that a competition with vitamin E could also take place with uptake transporters in the intestine and would thus lead to decreased vitamin K absorption.

The objectives of this study were to investigate the mechanisms of vitamin K\textsubscript{i} intestinal absorption, to evaluate the possibility of competition for absorption with vitamin E, and to specify the involvement of two transporters with broad specificity in this process: SR-BI\textsuperscript{2} and CD36.

MATERIALS AND METHODS

Chemicals

Phylloquinone (≥96% pure), 2-oleoyl-1-palmitoyl-sn-glycero-3-phosphocholine (phosphatidylcholine), 1-palmitoyl-sn-glycero-3-phosphocholine (lyso phosphatidylcholine), monoolein, free cholesterol, oleic acid, sodium taurocholate, dicholhexylcarbodiimide, hydroxy sulfosuccinimide sodium salt, and dimethylsulfoxide (DMSO) were purchased from Sigma-Aldrich. Vitamin K\textsubscript{1} 2,3-epoxide was obtained from Santa Cruz Biotechnology, Inc. R.R,R-α-Tocopherol (vitamin E) and echinenone (≥95% pure) were a generous gift of DSM Ltd. (Basel, Switzerland).

Mouse monoclonal IgG raised against the external domain (amino acids 104–294) of human SR-BI, also known as CLA-1, was purchased from BD Transduction Laboratories (Lexington, KY). Anti-human CD13 antibody (used as a control antibody against another plasma membrane protein) was from Santa Cruz Biotechnology. Mouse monoclonal anti-human CD36 IgM was purchased from Santa Cruz Biotechnology, Inc. R.R,R-α-Tocopherol (vitamin E) and echinenone (≥95% pure) were a generous gift of DSM Ltd. (Basel, Switzerland).

Preparation of Vitamin K\textsubscript{i}-enriched Vehicles for Cell and Mouse Experiments

Vitamin K\textsubscript{i}-rich Micelles—For delivery of phylloquinone to Caco-2 cells or to mouse intestinal explants, mixed micelles with lipid composition similar to those found in vivo (14) were prepared as described previously (15) to obtain the following final concentrations: 0.04 mm phosphatidylcholine, 0.16 mm lysophosphatidylcholine, 0.3 mm monoolein, 0.1 mm free cholesterol, 0.5 mm oleic acid, and 5 mm taurocholate (16). Phylloquinone was added into the micelles at a concentration of 0.25–5 μM, depending on the experiment. Concentration of vitamin K\textsubscript{i} in the micellar solutions was confirmed before each experiment.

Vitamin K\textsubscript{i}-rich Complete Medium—For delivery of phylloquinone to HEK cells, an appropriate volume of vitamin K\textsubscript{i} stock solution in ethanol was added to a glass tube to obtain a final concentration of 2.5 μM unless otherwise indicated. Stock solution solvent was carefully evaporated under nitrogen. Dried residue was solubilized into FBS overnight, and DMEM was then added to reach a final FBS concentration of 10%. The concentration of phylloquinone in the medium was confirmed before each experiment.

Vitamin K\textsubscript{i}-rich Emulsions—For delivery of phylloquinone to mice, emulsions were prepared as follows. An appropriate volume of stock solution containing 500 μg of phylloquinone was transferred to an Eppendorf tube. Stock solution solvent was carefully evaporated under nitrogen. Dried residue was solubilized in 100 μl of peanut oil (Lesieur, Asnières-sur-Seine, France), and 200 μl of a NaCl 0.9% solution was added. The mixture was vigorously mixed in an ice-cold water bath during sonication (Branson 3510) for 15 min and used for force-feeding within 10 min of preparation.

Cell Culture

Caco-2 Cell Culture—Caco-2 clone TC-7 cells (17, 18) were cultured in the presence of DMEM supplemented with 20% heat-inactivated FBS, 1% non-essential amino acids, and 1% antibiotics (complete medium), as described previously (13, 19). For each experiment, cells were seeded and grown on transwell plates for 21 days to obtain confluent and highly differentiated cell monolayers. Twelve hours prior to each experiment, serum-free complete medium was used in the apical and basolateral chambers.

HEK Cell Culture and Transfection—HEK 293-T cells were purchased from the American Type Culture Collection (Manassas, VA). Cells were cultured in 10% FBS complete medium at 37 °C in a humidified atmosphere of air/carbon dioxide (90:10, v/v) and the medium was changed every 48 h. Monolayers were subcultured with a 4-day passage frequency when they reached a confluence of about 80% and were subsequently treated with 0.25% trypsin-EDTA. For each experiment, cells were seeded at a 1:10 dilution in 6-well plates and transfected 24 h afterward with 3 μg of DNA and 6 μl of Jet Peptide in a 150 mm NaCl solution, according to the manufacturer’s instructions. The medium was then changed 10–12 h after, and cells were grown for an additional 24 h. Transfection efficiency was verified by Western blotting as described previ-

2 The abbreviations used are: SR-BI, scavenger receptor class B type I; CD36, cluster determinant 36; SR-BI transgenic mice, mice overexpressing SR-BI in the intestine; APCI, atmospheric pressure chemical ionization; SSO, sulfo-N-succinimidyl oleate; UPLC, ultraperformance liquid chromatography; HEK, human embryonic kidney.
Characterization of Vitamin \( K_1 \) Uptake in Cells

**Uptake Measurement**—At the beginning of each experiment, cell monolayers were washed with 0.5 ml of PBS. For uptake experiments, the apical side of the cell monolayers received the vitamin \( K_1 \)-enriched vehicles (whereas the other side received the serum-free complete medium if applicable). Cells were incubated for 30 min at 37 °C. At the end of each experiment, medium was harvested. Cells were washed twice in 0.5 ml of ice-cold PBS to eliminate adsorbed vitamin \( K_1 \) and then scraped and collected in 0.5 ml of ice-cold PBS. Absorbed vitamin \( K_1 \) was estimated as vitamin \( K_1 \) found in scraped cells plus vitamin \( K_1 \) found on the opposite side of the cell monolayer (if applicable).

**Apical Efflux Measurement**—For apical efflux experiments, the cells first received the vitamin \( K_1 \)-rich micelles at the apical side for 4 h. They were then washed three times with PBS and received apical medium containing vitamin \( K_1 \)-free mixed micelles (vitamin \( K_1 \) acceptors). At the end of each experiment, the apical medium was collected. Cells were washed twice in 0.5 ml of ice-cold PBS to eliminate adsorbed vitamin \( K_1 \) and then scraped and collected in 0.5 ml of PBS. Vitamin \( K_1 \) effluxed by cells (i.e., phylloquinone recovered in the apical medium) was expressed as a percentage of vitamin \( K_1 \) initially absorbed (determined as the total amount of vitamin \( K_1 \) recovered in cells, apical medium, and basolateral medium together if applicable).

**Competition with \( R,R,R \)-α-Tocopherol**—Phylloquinone uptake was measured after incubation with phylloquinone-rich mixed micelles \( (2.5 \, \mu M) \) containing α-tocopherol at concentrations ranging from 0 to 50 \( \mu M \). Similarly, α-tocopherol uptake was measured after incubation with tocopherol-rich micelles \( (5 \, \mu M) \) containing either no vitamin \( K_1 \) or 2.5 \( \mu M \) vitamin \( K_1 \).

**Transport Inhibition by BLT1**—For uptake experiments, BLT1 was used as described previously (13, 15, 21). Briefly, Caco-2 but not HEK cell monolayers were pretreated with either DMSO (control) or BLT1 at either 50 \( nM \) or 10 \( \mu M \) for 1 h. Both Caco-2 and HEK cells then received phylloquinone-enriched vehicles with either DMSO, 50 \( nM \) BLT1 or 10 \( \mu M \) BLT1, and uptake was measured as described above.

**Uptake Inhibition by Anti-human SR-BI Antibody**—For uptake experiments, cell monolayers were incubated for 5 min with 3.75 \( \mu g/mL \) anti-human SR-BI monoclonal antibody raised against the external domain before vitamin \( K_1 \)-rich micelles were added. Previous experiments have shown that this antibody concentration maximally inhibited absorption (15). Anti-human CD13 antibody was used as a control at 3.75 \( \mu g/mL \).

**Uptake Inhibition by SSO**—SSO can be used in concentrations up to 1.25 \( \mu M \), but it is classically used between 200 and 500 \( \mu M \) (13). Thus, during experiments, HEK cells received phylloquinone-enriched vehicles supplemented with 400 \( \mu M \) SSO, and uptake was measured as described above.

**Efflux Inhibition by BLT1**—The cells first received the phylloquinone-rich micelles at the apical side for 4 h. They were then washed two times with PBS and equilibrated in serum-free complete medium for 1 h. The cells then received apical medium containing vitamin \( K_1 \) acceptors (i.e., vitamin K-free micelles) supplemented with either DMSO or BLT1 at 10 \( \mu M \) for 3 h.

**Characterization of Vitamin \( K_1 \) Uptake in Mouse Intestine**

**Animals**—Wild-type (WT) mice, mice deficient in CD36 (22, 23), and mice overexpressing SR-BI in the intestine (24, 25) were housed in a controlled environment (temperature = 22 ± 2 °C, humidity = 55 ± 10%, darkness from 6 p.m. to 6 a.m.). They were given a standard A04 chow diet (Safe, Augy, France) and water ad libitum. They were fasted overnight before each experiment. The protocol (number 24-25092012) was approved by the local animal research ethics committee (Comité National de Réflexion Ethique sur l’Expérimentation Animale de Marseille 14).

**Vitamin \( K_1 \) Uptake Inhibition in Mouse Intestinal Explants**—Mice were fasted overnight, and intestinal explants were prepared as described previously (13). Briefly, on the day of the experiment, animals were euthanized by cervical dislocation, and the 4 cm distal to the first 3 cm of their small intestinal tract were immediately removed. The intestine samples were carefully rinsed with a sterile NaCl solution (0.9%), sliced into strips on ice as described previously (13), and homogeneously distributed in 12-well plates (Falcon Plates, BD Biosciences). Explants were incubated for 3 h in 500 \( \mu L \) of 2.5 \( \mu M \) vitamin \( K_1 \)-enriched mixed micelles supplemented with 10 \( \mu M \) BLT1 or 400 \( \mu M \) SSO or with no supplement. Each condition was performed in triplicate. After incubation, media were harvested, and the intestine explants were carefully rinsed twice in PBS. Samples were then suspended in 500 \( \mu L \) of PBS, homogenized with two 3-mm diameter stainless steel balls in 2-ml Eppendorf tubes using an MM301 ball mill (Retsch, Eragny sur Oise, France), and stored until analysis.

**Postprandial Experiment with Vitamin \( K_1 \)**—On the day of the experiment, the mice were force-fed with a phylloquinone-enriched emulsion.

To assess the phylloquinone postprandial response curve in WT mice, blood samples were taken by retroorbital puncture at 1.5, 3, 4.5, and 6 h after gavage. A final blood sample was taken at 7.5 h by intracardiac puncture before euthanasia by cervical dislocation.

To compare phylloquinone postprandial response in WT and transgenic mice, a blood sample was taken after 4 h of digestion by intracardiac puncture before euthanasia by cervical dislocation. The intestine of each animal was then quickly harvested, carefully rinsed with PBS, and cut in 6-cm segments (duodenum, proximal jejunum, median jejunum, distal jejunum, and ileum). All of the fragments were suspended in 500 \( \mu L \) of PBS and homogenized as described above before storage and analysis.

All of the samples were sealed under nitrogen and stored at −80 °C until vitamin \( K_1 \) analysis. Aliquots of cell samples were...
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used to assess protein concentrations using a bicinchoninic acid kit (Pierce).

Sample Acid Hydrolysis
To analyze the potential metabolism of phylloquinone to menadione in enterocytes, 500 μl of cell or WT intestinal sample (cf. above) were deconjugated via acid hydrolysis, which also oxidized the released menadiol to menadione, as described previously (26).

Vitamin K1 Extraction
Phylloquinone was extracted from 500 μl of aqueous sample using the following method. Distilled water was added to sample volumes below 500 μl to reach a final volume of 500 μl. Echinone was used as an internal standard and added to the samples in 500 μl of ethanol. The mixture was extracted once with 2 ml of hexane. The hexane phase obtained after centrifugation (500 × g, 10 min, 4 °C) was evaporated to dryness under nitrogen, and the dried residue was dissolved in 200 μl of mobile phase. A volume of 100–180 μl was used for HPLC analysis.

Vitamin K UPLC-MS Analysis
Analyses were performed by ultraperformance liquid chromatography using an ACQUITY system (Waters Corp., Milford, MA) linked simultaneously to a diode array detector (UPLC DAD, Waters Corp.) and interfaced with a Bruker Daltonics HCT Ultra Ion Trap mass spectrometer via an atmospheric pressure chemical ionization source operated in positive ion mode (UPLC DAD/APCI-MS). Compass™ software (Bruker Daltonics, Bremen, Germany) was used for mass spectrometric instrument control and data processing.

UPLC conditions were modified from Song et al. (27). Separation was carried out using an ACQUITY UPLC® BEH C18 column (50 mm × 2.1 mm, inner diameter 1.7 μm; Waters). The column temperature was 35 °C, the flow rate was 0.45 ml min⁻¹, and 10 μl was injected. Analysis was achieved using a gradient of Milli-Q water/2-propanol (80:20, v/v) containing 0.1% HCOOH (v/v) (solvent A) and CH₃OH containing 0.1% HCOOH (v/v) (solvent B). Gradient conditions of the mobile phase were as follows: A, 85%, 0–3 min; A, 85–5%, 3–8 min; A, 2%, 8–12 min; A, 2–0%, 12–14 min, followed by a return to the initial conditions (solvent A, 85%) and re-equilibration for 4 min. Peaks were detected at 270 nm.

The ion trap was operated in the Ultra Scan mode from m/z 120 to 800. The ICC target was set to 100,000 with a maximum accumulation time of 200 ms. Nitrogen (99.99% purity) was used as the desolvation gas. The source parameters were set as follows: dry temperature 350 °C, nebulizer pressure 50 p.s.i., dry gas flow 5 liters/min⁻¹, vaporizer temperature of 350 °C, capillary voltage −2 kV, and corona current 4000 nA.

Vitamin K HPLC Analysis
Phylloquinone and RRR-α-tocopherol were separated using a 250 × 4.6-nm RP C₁₈, 5-μm Zorbax column (Interchrom, Montluçon, France) and a guard column.

The isocratic mobile phase consisted of 80% methanol, 19.45% ethanol, and 0.55% H₂O containing 1 mm sodium acetate, 2 mm zinc chloride, and 2 mm acetic acid (flow rate = 1.2 ml/min⁻¹), and the column was kept at a constant temperature (35 °C). The HPLC system was composed of a Dionex separation module (P680 HPLC pump and ASI-100 automated sample injector, Dionex, Aix-en-Provence, France), a Dionex UVD340U photodiode array detector, and a Jasco fluorimetric detector (Jasco, Nantes, France). Depending on the quantity of vitamin K present in the sample, phylloquinone was either measured at 248 nm using the photodiode array detector or at 430 nm after light emission at 244 nm using fluorimetric analysis (after postcolumn online reduction to the hydroquinone form in a 4.6 mm × 5-cm column freshly filled with 98% pure zinc dust) and identified by retention time coincident with authentic standard. Quantification was performed using Chromeleon software (version 6.50 SP4 Build 1000) comparing peak area with external calibration curves.

Statistical Analysis
Results are expressed as means ± S.E. Differences between more than two groups of unpaired data were analyzed using the non-parametric Kruskal-Wallis test. The non-parametric Mann-Whitney test was used as a post hoc test when the Kruskal-Wallis test showed significant differences between groups. Differences between only two groups of unpaired data were tested via Mann-Whitney U test. Values of p < 0.05 were considered significant. All statistical analyses were performed using Statview software, version 5.0 (SAS Institute, Cary, NC).

RESULTS

Metabolism of Vitamin K₁ by Caco-2 Cells and Mouse Enterocyte—We first aimed to assess whether phylloquinone was metabolized in Caco-2 cells and mouse intestine.

Using the UPLC DAD/APCI-MS conditions described above, vitamin K₁ was detected at 11.8 min (Fig. 1A, compound 2), producing a protonated ion [M + H]+ with m/z 451.2. After fragmentation, a daughter ion was detected at m/z 186.8 (Fig. 1B). The analyses of the cell samples incubated with vitamin K₁ for either 1 or 24 h did not reveal local production of menadione (data not shown), regardless of whether or not the samples were acid-hydrolyzed. No other metabolites were detected after a 1-h incubation. However, vitamin K₁ epoxide was detected after 24 h of incubation (Fig. 1A, compound 1) in the cellular compartment only of both acid hydrolyzed and non-hydrolyzed samples (retention time 10.8 min, m/z 467.2, identical to the standard molecule) (Fig. 1C). To avoid the metabolism of phylloquinone by the cells, cell experiments were conducted with incubation periods of 1 h. UPLC-MS analyses of WT mouse intestinal samples 4 h after force feeding demonstrated that vitamin K₁ was the only form of vitamin K detectable in the intestinal mucosa in our experimental conditions (data not shown).

Effect of Micellar Vitamin K₁ Concentration on Vitamin K₁ Uptake by Caco-2 Cells—We then aimed to characterize vitamin K₁ uptake by differentiated Caco-2 cell monolayers.
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At concentrations ranging from 0.5 to 5 μM, absorption of vitamin K₁ was not linear (Fig. 2A). The best fit was achieved using a hyperbolic curve \( y = ax/(x + b) \), \( R^2 > 0.99 \). Apparent \( Q_{\text{max}} \) was 1.47 nmol of phylloquinone/mg of protein, and \( K \) was equal to 8.58 μM. \( Q_{\text{max}} \) represents the maximal amount of phylloquinone absorbed, and \( K \) represents the concentration of micellar phylloquinone required to reach \( Q_{\text{max}}/2 \).

Effect of Vitamin E Co-incubation on Vitamin K₁ Uptake by Caco-2 Cells and Vice Versa—Next, we evaluated whether competition between vitamin K₁ and E exists during nutrient uptake in differentiated Caco-2 cell monolayers. As shown in Fig. 2B, co-incubation of 5 μM micellar \( \alpha \)-tocopherol (vitamin E) with 2.5 μM micellar phylloquinone significantly decreased the uptake of phylloquinone by 28.0% (\( p < 0.05 \)). Similarly, co-incubation of 2.5 μM micellar phylloquinone with 5 μM micellar \( \alpha \)-tocopherol significantly decreased the uptake of \( \alpha \)-tocopherol by 29.2% (\( p < 0.05 \)). This effect was dose-dependent, as further shown in Fig. 2C, which suggests that these two vitamins share common uptake pathways.

Effect of the Amount of Cholesterol Incorporated in Micelles with Vitamin K₁ on Vitamin K₁ Uptake—Cholesterol is naturally present in micelles during digestion, but its concentration may vary depending on diet and biliary secretion. Fig. 2D shows that high micellar cholesterol concentrations significantly affected phylloquinone uptake. This suggests that cholesterol and phylloquinone may also display common uptake pathways.

Interestingly, vitamin E and cholesterol share common protein carriers at the apical side of the enterocyte. Among these transporters, the scavenger receptors SR-BI and CD36 are known to display broad substrate specificity. We thus investigated the involvement of SR-BI and CD36 in vitamin K₁ intestinal apical uptake.

Involvement of SR-BI in Vitamin K₁ Transport across the Apical Membrane of Caco-2 Cells

Uptake Experiment—The specific chemical inhibitor of SR-BI, BLT1, significantly decreased apical Caco-2 cell phylloquinone uptake by up to 38.5% when added at 50 nM (~IC₅₀ (28)). When added in excess (10 μM (13, 15, 21, 25, 29)), it led to an inhibition of 84.5% of phylloquinone uptake (Fig. 3A, \( p < 0.05 \)). Moreover, Fig. 3B shows that the addition of anti-human SR-BI antibody raised against the external domain significantly decreased phylloquinone apical uptake by 15.6% compared with the control antibody (anti-CD13 antibody, \( p < 0.05 \)). CD36 involvement in vitamin K₁ uptake was not challenged in Caco-2 cells because this cell line does not express the protein.

Apical Efflux Experiment—Caco-2 cells were challenged in phylloquinone before apical efflux of vitamin K₁ was measured in the presence or the absence of BLT1 (Fig. 3C). Adding 10 μM BLT1 to the apical side of the cells in the presence of vitamin K-free mixed micelles significantly reduced the apical efflux of vitamin K₁ by 80.5% (Fig. 2B, \( p < 0.05 \)).

Effect of SR-BI and CD36 Transfection on Vitamin K₁ Uptake in HEK Cells—To validate our previous and to extend then to CD36, we then used transfected HEK cells. Transfection with human SR-BI led to a significant 109.6% increase of phylloquinone uptake compared with control (cells transfected with an empty plasmid), and this increase was totally suppressed by 10 μM BLT1 (Fig. 4A). Fig. 4B shows that HEK cell transfection with human CD36 significantly increased phylloquinone uptake by 164.3% and that this increase was significantly impaired by the addition of a chemical inhibitor of CD36, SSO (400 μM). Because SR-BI and CD36 are not endogenously expressed in HEK cells (Fig. 3, insets), neither BLT1 nor SSO had an effect on phylloquinone uptake in untransfected HEK cells (data not shown).

Effect of BLT and SSO on Vitamin K₁ Uptake by Mouse Intestinal Explants—To further confirm our data, we next performed ex vivo experiments. BLT1 and SSO significantly impaired phylloquinone uptake by mouse intestinal explants (~19.4 and ~18.9%, respectively; Fig. 5).

Vitamin K₁ Postprandial Response in Mice—Finally, we assessed the phylloquinone postprandial response in mice after gavage (Fig. 6). Each mouse received 500 μg of phylloquinone in an oil-in-water emulsion. Our data show that phylloquinone peaked in plasma between 1.5 and 4.5 h after gavage.

Effect of Intestinal SR-BI Overexpression on Vitamin K₁ Absorption in Mice—After gavage, the phylloquinone postprandial response was significantly increased in SR-BI transgenic mice compared with control mice (±162.4%, \( p = 0.0071 \); Fig. 6A). This difference was still significant after adjustment by plasma total lipids (±548.9%, \( p = 0.0011 \); Fig. 7A). Further-
more, the phylloquinone content of the first two fragments (duodenum and proximal jejunum) of SR-BI transgenic mice was significantly higher than the vitamin K\(_1\) content of the first two fragments of WT mice (+302.4%, \(p = 0.0008\) and +183.1%, \(p = 0.0116\); Fig. 7B). Conversely, the phylloquinone content of the last three fragments (median jejunum, distal jejunum, and ileum) of SR-BI transgenic mice was lower than the vitamin K\(_1\) content of the matched fragments of WT mice, although the difference was significant in the distal jejunum only (-69.1%, \(p = 0.0324\); Fig. 5B).

**Effect of CD36 Deletion on Vitamin K\(_1\) Absorption in Mice**—After gavage, the phylloquinone postprandial response was significantly increased in CD36-deficient mice compared with control mice (+290.2%, \(p = 0.014\); Fig. 7C). This difference did not remain significant after adjustment by plasma total lipids. Additionally, the phylloquinone content in all intestinal fragments remained similar in the two groups of mice (Fig. 7D).

**DISCUSSION**

Phylloquinone intestinal absorption has been poorly investigated so far. To evaluate in detail the mechanism(s) involved in transport across the enterocyte, we used the popular Caco-2 TC-7 cell model, which has been extensively used to study fat-soluble vitamin intestinal absorption (13, 21, 30). There are no data on phylloquinone micellar concentration in the human gut. However, previous work has shown that lutein concentrations in the human duodenal lumen were 5–7 \(\mu M\) after a spinach-rich meal providing \(\sim 10\) mg lutein (31). Considering spinach phylloquinone content (32), such a meal would result in the production of micelles containing \(\sim 0.03–0.05 \mu M\) phylloquinone. However, to allow an accurate quantification in cells, we used micellar concentrations ranging from 0.25 to 5 \(\mu M\) and principally 2.5 \(\mu M\).

We first aimed to assess whether vitamin K\(_1\) was metabolized by intestinal cells under our experimental conditions. APCI is the most common method of ionization for lipophilic compounds in general, including vitamin K (27, 33, 34). The best detection was obtained in positive mode; thus, this was used for our analysis. The \(m/z\) 186.9 daughter ion corresponds to the loss of the side chain, as reported previously (34). A vitamin K\(_1\) epoxide was identified in the cytosol of Caco-2 cells after a 24-h incubation of vitamin K\(_1\)-rich mixed micelles, whereas no metabolites were detected after a 1-h incubation. To avoid phylloquinone degradation and metabolism to accurately measure vitamin K\(_1\) transport, all experiments in Caco-2 cells were conducted after short incubation periods, and samples were analyzed by HPLC.

Our first results showed that vitamin K\(_1\) uptake is saturable, which supports a protein-mediated uptake hypothesis (35). Apparent \(Q_{\text{max}}\) for phylloquinone was in the same range as the apparent \(Q_{\text{max}}\) for tocopherol (21). The apparent \(K\) for phyllo-
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Phylloquinone was 10-fold smaller than that of tocopherol, suggesting a greater affinity of the cell brush-border membrane for vitamin K₁ than for vitamin E, but these data should be considered cautiously because these values strongly depend on both micelle composition and experimental conditions (Caco-2 cells differentiated on filters, incubation period, etc.). The saturable uptake of vitamin K₁ by Caco-2 cells is consistent with the previous findings of Holland and colleagues, who showed that the use of a nitrogen atmosphere to inhibit mitochondrial respiration and the addition of 2,4-dinitrophenol to inhibit ATP synthesis together impaired phylloquinone intestinal absorption in rats. Although nonspecific experiments, these data suggested the involvement of a carrier protein in phylloquinone uptake (36).

We then explored the possibility of competition between dietary vitamin K₁ and α-tocopherol for absorption. The fact that both vitamins could impair the uptake of each other suggests that the two molecules compete for entry into the intestinal cell and thus may share common absorption pathway(s). This situation could be particularly deleterious for vitamin K because its dietary intake is moderate (recommended daily allowance for an adult = 75 μg/day) compared with vitamin E (recommended daily allowance of 15 mg/day), with vitamin E supplementation increasing the imbalance further. This result could at least partially explain the excessive bleeding observed in patients supplemented with vitamin E (11). This result seems in disagreement with previous data obtained in rats and showing a non-significant decrease of phylloquinone serum concentration 6 h after gavage with 0.2 mg of phylloquinone and 10 mg of α-tocopherol (12). However, our data suggest that the phylloquinone absorption peak is probably earlier, around 3–4 h (Fig. 6), and a single measure of phylloquinone postprandial concentration at 6 h cannot exclude possible competition for absorption.

Vitamin E intestinal absorption was shown to involve cholesterol transporters, including scavenger receptors (21, 37). We thus aimed to evaluate the involvement of SR-BI and CD36 in...
phylloquinone transport across the brush border membrane. The finding that phylloquinone uptake in Caco-2 cells was specifically inhibited by both anti-human SR-BI antibody and BLT1 (a chemical inhibitor of SR-BI) (38, 39), is taken as primary evidence that SR-BI is involved in vitamin K₁ uptake. BLT1 led to a 80% decrease in phylloquinone uptake, whereas the antibody led to a 15% decrease only. This difference, already observed previously with vitamin E (21), can be explained by the fact that BLT1 may be more efficient than the antibody to inhibit SR-BI activity. Moreover, the presence of proteases at the brush border membrane may hydrolyze the antibodies. Interestingly, the role of SR-BI is not restricted to vitamin K₁ uptake alone. The transporter was also able to mediate vitamin K₁ efflux from the cell to the apical medium, as has been similarly shown for vitamin E (21) and D (13). CD36 involvement could not be evaluated in the Caco-2 monolayers because this cell model does not express the protein (15). Thus, in order to further examine the involvement of both scavenger receptors in phylloquinone transport, we studied phylloquinone uptake using HEK cells transfected with either SR-BI or CD36. HEK cells transfected with SR-BI displayed an increased BLT1-sensitive phylloquinone uptake, confirming SR-BI involvement in vitamin K₁ absorption. Similarly, HEK cells transfected with CD36 exhibited SSO-sensitive uptake of phylloquinone, suggesting the involvement of CD36. We then performed an ex vivo experiment using mouse intestinal explants. We observed a significant 20% inhibition of vitamin K₁ uptake by BLT1. This inhibition was less pronounced than the inhibition observed in Caco-2 cells treated with BLT1 (80%). This discrepancy is probably due to different transporter expression profiles in vivo and in vitro. We also confirmed the results obtained in HEK cells because SSO impaired phylloquinone uptake by mouse intestinal mucosa. However, the fact that we did not manage to fully inhibit phylloquinone uptake suggests that passive diffusion or other transporters are, at least partially, involved.

Finally, we performed in vivo absorption experiments. Preliminary tests showed that vitamin K₁ was not detectable in fasting plasma samples of mice bred in house, and no vitamin K metabolites were detected in mouse intestinal mucosa 4 h after force feeding. The results obtained in mice overexpressing SR-BI...
in the intestine provide further evidence that SR-BI is directly involved in phylloquinone absorption. Indeed, the phylloquinone postprandial response was significantly increased in transgenic mice compared with wild-type animals. Furthermore, phylloquinone content of the intestinal mucosa of the transgenic mice was very different from the controls. Phylloquinone significantly accumulated in the first fragments of mouse intestine, where SR-BI overexpression is the strongest (24). Conversely, less phylloquinone uptake was recovered in the last intestinal sections, probably because it was largely absorbed in the proximal part of the intestine.

The results obtained in CD36-deficient mice revealed an indirect involvement of this transporter in phylloquinone transport. Indeed, CD36-deficient mice had a higher phylloquinone postprandial response than wild-type mice, contrary to what might be expected if CD36 is mainly implicated in phylloquinone uptake. Deficient mice also had a significantly increased postprandial triglyceride response (data not shown). This result is in agreement with our previous work on γ-tocopherol (37). This result is also in agreement with previous data demonstrating that in the presence of lipids, CD36 induces large chylomicron formation via an induction of both apolipoprotein B48 and microsomal triglyceride transfer protein, probably through ERK1/2 modulation. By contrast, CD36 deletion results in an impaired secretion of chylomicrons that are smaller in size and thus resistant to lipoprotein lipase activity (20, 23, 40), which leads to triacylglycerol and phylloquinone accumulation. The fact that phylloquinone content was similar in WT and CD36-deficient mouse intestinal fragments indicates that CD36 involvement in vitamin K uptake may be balanced by the activity of other transporters. Another possibility is that CD36 also impacts chylomicron secretion from the enterocyte to the lymph. A delay in secretion would then result in a temporary accumulation of the chylomicrons in the intestinal cells that would mask a reduced uptake. Because our cell models (CD36-transfected HEK cells) cannot mimic this secretion step, this could also explain the discrepancy between our in vitro and in vivo data. Further studies are required to validate this last assumption.

The physiological consequences of these novel findings remain to be evaluated. We suggest that interindividual variation in intestinal scavenger receptor expression and/or function may be linked to variations in intestinal phylloquinone absorption efficiency. Ultimately, these variations could impact health status in subgroups of population at risk of developing cardiovascular diseases or osteoporosis.

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