

Inhibition of ERK1/2 improves lipid balance in rat macrophages via ABCA1/G1 and CD36

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Abstract. ATP-binding cassette transporters A1 (ABCA1) and G1 (ABCG1), and macrophage scavenger receptor, cluster of differentiation (CD)36, function as key mediators of cholesterol efflux and influx from macrophages. In addition, they are associated with foam cell formation and the development of atherosclerosis (AS). The aim of the present study was to investigate the effects of extracellular signal-regulated kinases 1/2 (ERK1/2) inhibition on lipid balance in oxidized-low-density lipoprotein (Ox-LDL)-stimulated rat macrophages, and to examine the role of ERK1/2 inhibitors in AS. Rat peritoneal macrophages were treated with Ox-LDL alone or in combination with an ERK1/2 inhibitor, U0126, and untreated cells served as controls. Ox-LDL-induced lipid accumulation was detected by DiI fluorescence and oil red O staining. In addition, the mRNA and protein expression levels of ABCA1, ABCG1 and CD36 were determined using polymerase chain reaction and western blotting, respectively. Treatment with Ox-LDL significantly increased lipid accumulation and upregulated

the mRNA and protein expression levels of ABCA1, ABCG1 and CD36 in macrophages. The addition of U0126 resulted in a marked reduction of lipid deposition, upregulation of ABCA1/G1 expression and suppression of CD36 expression in Ox-LDL-stimulated macrophages. The results of the present study indicated a novel association between ERK1/2 signaling and lipid metabolism, thus suggesting that inhibition of ERK1/2 may be considered a promising therapeutic strategy against AS.

Introduction

Atherosclerosis (AS) is a progressive disease characterized by accumulation of lipids and fibrous elements in the arterial tunica intima, which results in atherosclerotic plaque formation and arterial narrowing (1). During the progression of AS, macrophages are critical (2). The development of AS is strongly associated with engulfment of oxidized-low-density lipoprotein (Ox-LDL) by macrophages. It has previously been reported that macrophages engulf Ox-LDL via scavenger receptors, resulting in the deposition of a large quantity of lipids, which further promote the progression of AS (1). Recently, mitogen-activated protein kinase (MAPK) signaling cascades have been investigated in macrophages *in vitro*, in order to evaluate their importance (3).

Extracellular signal-regulated kinases 1/2 (ERK1/2) are members of the MAPK family, which are involved in mediating various processes, including cell cycle progression, cell migration, survival, differentiation and proliferation (4). However, the association between ERK1/2 signaling and lipid metabolism remains to be elucidated.

ERK1/2 inhibitors have been reported to synergize with liver X receptor (LXR) ligand to induce ATP-binding cassette transporter (ABC)A1 expression and regulate cholesterol efflux (5). In addition, ABCA1 and ABCG1 have been demonstrated to stimulate the efflux of lipids via the reverse cholesterol transport (RCT) pathway (6). As a specific macrophage scavenger receptor, cluster of differentiation (CD)36 recognizes and internalizes Ox-LDL particles, resulting in cellular cholesterol uptake, lipid accumulation in intracellular space and the production of foam cells (7). It has therefore been hypothesized that ABCA1/G1 and CD36

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Abbreviations: AS, atherosclerosis; RPMs, rat peritoneal macrophages; Ox-LDL, oxidized-low-density lipoprotein; DiI-ox-LDL, DiI-labeled oxidized-low-density lipoprotein; ABCA1, ATP-binding cassette transporter A-1; ABCG1, ATP-binding cassette transporter G-1; MAPK, mitogen-activated protein kinase; ERK1/2, extracellular signal-regulated kinases 1/2; p-ERK, phosphorylated ERK; PBS, phosphate-buffered saline

Key words: extracellular signal-regulated kinases 1/2, macrophage, lipid metabolism, ATP-binding cassette transporter A-1, CD36, ATP-binding cassette transporter G-1

are crucial to the maintenance of intracellular cholesterol stability or the efficacy of the RCT process (8). However, the regulation of ABCA1/G1 and CD36 expression by ERK1/2 in macrophages remains to be elucidated. The aim of the present study was to investigate the association between ERK1/2 and lipid metabolism, and evaluate the effect of ERK1/2 on the expression levels of ABCA1/G1 and CD36 in macrophages.

Materials and methods

Animals. A total of 40 male Sprague Dawley rats (weight, 200–220 g; age, 8–9 weeks) were purchased from the Laboratory Animal Center of Fujian Medical University (Fuzhou, China). The rats were housed under a 12 h light/dark cycle at 37°C in an atmosphere containing 5% CO₂, and were given *ad libitum* access to food and water. All animal experiments in the present study were performed in accordance with the recommendations of the Guidelines for the Care and Use of Laboratory Animals of the Ministry of Science of People's Republic of China. The present study was approved by the Animal Care and Use Committee and Institutional Review Board of Fujian University of Traditional Chinese Medicine (Fuzhou, China) (permission no. IRB201100117) and the Ethics Review Committee of Fujian University of Traditional Chinese Medicine (permission no. FJZYDXERC2011010).

Isolation and treatment of rat peritoneal macrophages (RPMs). The rats were anesthetized with 10% chloral hydrate (Peking Union-Biology Co., Ltd., Beijing, China) and were sacrificed by cervical dislocation, and soaked in 75% ethanol for 5 min. RPMs were harvested by lavaging the peritoneal cavity with 10 ml Dulbecco's modified Eagle's medium/Ham's F12 (DMEM-F12; Hyclone; GE Healthcare Life Sciences, Logan, UT, USA). Cells were centrifuged at 4°C, 1,000 × g for 15 min, and cultured in DMEM supplemented with 1% glutamine and penicillin/streptomycin (Gibco; Thermo Fisher Scientific, Inc., Waltham, MA, USA), and 10% fetal bovine serum (FBS; Hyclone; GE Healthcare Life Sciences) at 37°C in an incubator containing 5% CO₂. RPMs in the exponential growth phase were seeded onto 6-well plates at a density of 5 × 10⁶ cells/well for the indicated time. RPMs were randomly assigned to three groups. Cells in the control group were incubated in DMEM-F12 supplemented with 10% FBS for 12, 24 and 48 h; cells in the Ox-LDL group were treated with 50 mg/l Ox-LDL (Peking Union-Biology Co., Ltd.) or 10 µg/ml DiI-Ox-LDL (Peking Union-Biology Co., Ltd.) for 12, 24 and 48 h, respectively (8–12); whereas cells in the Ox-LDL + U0126 group were incubated in DMEM-F12 containing 10% FBS supplemented with 10 µM U0126 (Sigma-Aldrich, St. Louis, MO, USA) and Ox-LDL or DiI-Ox-LDL for 12, 24 or 48 h.

Morphological observation. RPMs were incubated with DMEM-F12 containing 10% FBS in 6-well plates. Once the cells had adhered to the culture plates, RPMs were pretreated with Ox-LDL alone, or in combination with U0126, for 12, 24 and 48 h, whereas the untreated cells served as controls. Cellular lipid accumulation was detected using oil red O staining and DiI fluorescence. Cells were washed three times with phosphate-buffered saline (PBS), fixed in 5% formalin

solution for 15 min at room temperature, washed once with PBS and stained with oil red O (Wuhan Boster Biological Technology, Ltd., Wuhan, China) for 10 min, followed by hematoxylin (Wuhan Boster Biological Technology, Ltd.) staining for 5 min. Cells were then observed under a Leica DM IL compact inverted stereo microscope (Leica Microsystems GmbH, Wetzlar, Germany), and images were captured at magnification x400. In addition, RPMs in monolayer cultures were exposed to DiI-Ox-LDL alone, or in combination with U0126, for 12, 24 and 48 h. The medium was removed and cells were washed with PBS, mounted on cover slips and analyzed using fluorescent microscopy.

Cytotoxicity assay. The viability of RPMs post-treatment with Ox-LDL (10, 30, 50, 70 and 90 mg/l) was assessed using an MTS assay (Promega Corporation, Madison, WI, USA). The RPMs were grown in 96-well plates and were incubated with Ox-LDL (10, 30, 50, 70 or 90 mg/l) for 12, 24 or 48 h. The MTS assay was used to measure the viability of the cells. Cells were incubated at 37°C with MTS (1.90 mg/ml) for 4 h, and absorbance was measured using a microplate reader (Multiskan GO; Thermo Fisher Scientific Inc.) at a wavelength of 490 nm. Three wells were set for each concentration of Ox-LDL. All experiments were repeated twice.

Reverse transcription-polymerase chain reaction (RT-PCR) assay. Total RNA was isolated from the RPMs using TRIzol® (Invitrogen; Thermo Fisher Scientific, Inc.), and total RNA (500 ng) was reverse transcribed in a total volume of 20 µl containing oligo (dT) primers at 42°C for 30 min using TransScript® One-Step gDNA Removal and cDNA Synthesis SuperMix (Beijing Transgen Biotech Co., Ltd., Beijing, China). Primers used for PCR (Thermo Fisher Scientific, Inc.) are presented in Table I. PCR was performed on an Applied Biosystems® 2720 Thermal Cycler (Applied Biosystems; Thermo Fisher Scientific, Inc.) in a 20 µl reaction system containing cDNA (500 ng/µl), each specific primer, and 2X EasyTaq® PCR SuperMix (Beijing Transgen Biotech Co., Ltd.) under the following conditions: Pre-denaturation at 94°C for 5 min; 35 cycles at 94°C for 30 sec, 58°C for 30 sec and 72°C for 1 min; followed by final extension at 72°C for 7 min. The PCR products were separated by 1.5% agarose gel electrophoresis, and the relative mRNA expression levels were determined as the ratio of the grayscale value of the target gene to GAPDH (Image-Lab version 5.0; Bio-Rad Laboratories, Inc., Hercules, CA, USA). The experiment was repeated three times.

Western blot analysis. RPMs were digested with pancreatin (Gibco; Thermo Fisher Scientific, Inc.), pipetted evenly, and centrifuged. The supernatant was transferred to a 1.5 ml centrifuge tube, lysed on ice in radioimmunoprecipitation assay lysis buffer (Beyotime Institute of Biotechnology, Shanghai, China) for 30 min, and fractionated three times with ultrasound. The mixture was then centrifuged at 13,800 × g for 15 min, and the supernatant was stored at -80°C for subsequent experiments. Protein isolated from the RPMs was quantified using the Bicinchoninic Acid assay (Wuhan Boster Biological Technology, Ltd). Approximately 30 µg protein was separated by 10% sodium dodecyl sulfate-polyacrylamide gel electrophoresis at 110 V for 2.5 h at room temperature, and

Table I. Sequences of primers used for reverse transcription-polymerase chain reaction.

Gene	Primer sequence, 5'→3'
CD36	F: ACTCCAGAACCCAGACAACCAC R: ACCAAGTAAGACCATCTCAACCAG
ABCA1	F: CCTAAGCATTATCAAGGAGGGAAG R: AGAGATGACAAGGAGGACGGAAG
ABCG1	F: GTCCTGGGCATCTTCTTCATCTC R: CCAACTCAGCCAACACTCCTCTC
GAPDH	F: ACGGCAAGTTCAACGGCACAG R: GAAGACGCCAGTAGACTCCACGAC

CD36, cluster of differentiation 36; ABCA1, ATP-binding cassette transporter A-1; ABCG1, ATP-binding cassette transporter G-1; GAPDH, glyceraldehyde 3-phosphate dehydrogenase; F, forward; R, reverse.

was then transferred to polyvinylidene difluoride membranes (EMD Millipore, Billerica, MA, USA) for a further 1.5 h at 110 V and 4°C. The membrane was blocked with 5% non-fat milk in Tris-buffered saline for 1 h at room temperature and probed with mouse monoclonal anti-ABCA1 (1:1,000; cat. no. ab18180), rabbit monoclonal anti-ABCG1 (1:1,000; cat. no. ab52617), rabbit polyclonal anti-CD36 (1:500; cat. no. ab78054) (Abcam, Cambridge, UK), rabbit polyclonal phosphorylated (p)-p44/42 MAPK (ERK1/2) (1:1,000; cat. no. 9101), rabbit polyclonal p44/42 MAPK (ERK1/2) (1:1,000; cat. no. 9102) and anti-β-actin (1:1,000; cat. no. 4967) (Cell Signaling Technology, Inc., Danvers, MA, USA) antibodies at 4°C overnight. The membrane was then washed with PBS, and incubated with horseradish peroxidase-conjugated goat anti-rabbit (1:5,000; cat. no. A0208) and anti-mouse (1:5,000; cat. no. A0216) secondary antibodies (Beyotime Institute of Biotechnology) for a further 2 h at room temperature. Enhanced chemiluminescence reagents (Beyotime Institute of Biotechnology) and chemiluminescent substrates were used to visualize immunoreactive bands. The grayscale values of the protein bands were estimated using the image processing software Image-Lab version 5.0 (Bio-Rad Laboratories, Inc.). The ratio of the grayscale value of the target protein band to β-actin protein band was defined as the protein expression level. All experiments were repeated three times.

Statistical analysis. All data are presented as the mean ± standard deviation. Differences between means were determined using one-way analysis of variance followed by Dunnett's test, or Student's t-test. Statistical analyses were conducted using SPSS version 18.0 (SPSS, Inc., Chicago, IL, USA). $P < 0.05$ was considered to indicate a statistically significant difference.

Results

Oil red O staining. RPMs were treated with 50 mg/l Ox-LDL for 12, 24 and 48 h, washed twice with PBS, and stained with oil red O and hematoxylin. Intense oil red O staining (red) was detected in RPMs treated with Ox-LDL, indicating intracellular

lipid accumulation. However, staining was markedly reduced in RPMs co-treated with Ox-LDL and 10 μM U0126, suggesting reduced lipid deposition (Fig. 1A).

Fluorescence. To examine the effects of the ERK1/2 inhibitor on DiI-Ox-LDL uptake by the RPMs, cells were cultured with DiI-Ox-LDL for 12, 24 and 48 h. Treatment with DiI-Ox-LDL markedly increased cytoplasmic fluorescence of RPMs, as compared with untreated controls (Fig. 1B), and engulfment of DiI-Ox-LDL in RPMs was demonstrated by observation of intracellular accumulation of DiI-labeled lipids. However, markedly reduced DiI-Ox-LDL fluorescence was observed in RPMs following the addition of U0126, indicating that treatment with U0126 may result in reduced lipid deposition.

MTS assay. To evaluate the hypothesis that Ox-LDL may result in necrosis and apoptosis of RPMs, the effect of Ox-LDL on cell viability was evaluated by MTS assay. A greater viability of RPMs was observed following treatment with Ox-LDL at a concentration of 50 mg/l, as compared with Ox-LDL treatment at 10, 30, 70 and 90 mg/l for 12 and 48 h. Treatment with Ox-LDL at a concentration of 50 mg/l resulted in greater viability of RPMs, as compared with Ox-LDL treatment at 70 and 90 mg/l for 24 h. These results suggest that cell viability was increased in response to 50 mg/l Ox-LDL treatment (Fig. 2A).

Expression levels of ERK and p-ERK. Western blot analysis detected comparative ERK expression in RPMs treated with Ox-LDL for 12, 24 and 48 h, whereas p-ERK expression levels were markedly increased in Ox-LDL-treated RPMs, as compared with untreated RPMs (Fig. 2B). The addition of 10 μM U0126 led to a marked reduction in the expression levels of p-ERK, as compared with treatment with Ox-LDL alone, indicating that the ERK1/2 inhibitor attenuates ERK1/2 phosphorylation.

Inhibition of ERK1/2 upregulates Ox-LDL-induced ABCA1 expression. To evaluate the effect of ERK1/2 inhibition on ABCA1 expression, mRNA and protein expression levels of ABCA1 were determined in Ox-LDL-treated RPMs. ABCA1 expression levels were markedly elevated in RPMs treated with Ox-LDL alone or Ox-LDL + U0126 at the mRNA (Fig. 3A) and protein (Fig. 3B) level. In addition, the mRNA and protein expression levels of ABCA1 were notably increased in RPMs following co-treatment with Ox-LDL + U0126 for 12 and 24 h; however, the mRNA and protein expression levels of ABCA1 were not significantly greater in RPMs treated with Ox-LDL + U0126 for 48 h, as compared with RPMs treated with Ox-LDL alone. These results indicate that ERK1/2 inhibition may upregulate mRNA and protein expression levels of ABCA1 in Ox-LDL-treated macrophages.

Inhibition of ERK1/2 augments Ox-LDL-induced ABCG1 expression. Western blotting and RT-PCR indicated a significant increase in ABCG1 expression at the transcriptional (Fig. 4A) and translational (Fig. 4B) level following Ox-LDL treatment. In addition, mRNA and protein expression levels of ABCG1 were significantly higher in RPMs co-treated with Ox-LDL and U0126 for 12 and 24 h, as compared with those treated with Ox-LDL alone. Conversely,

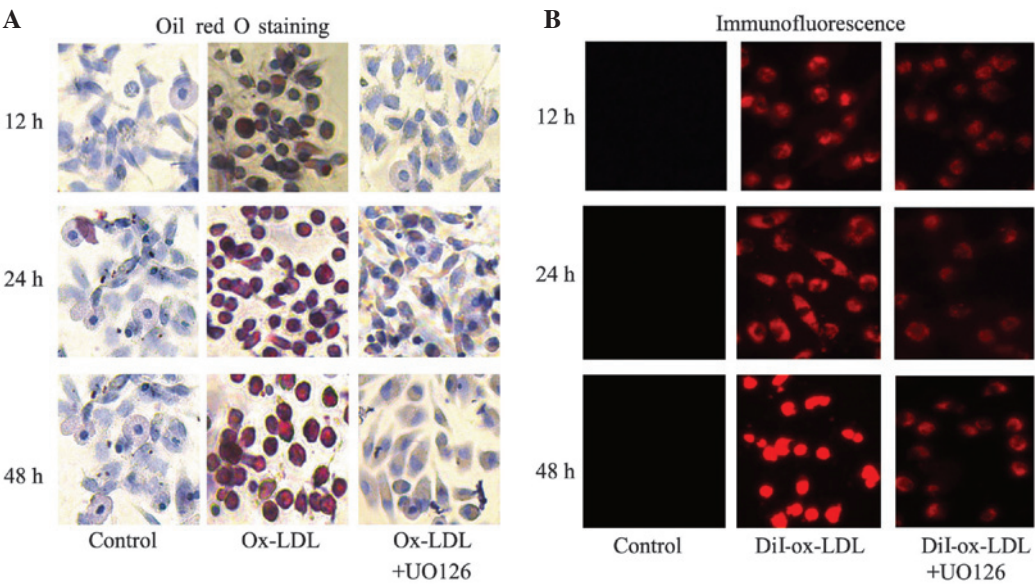


Figure 1. ERK1/2 inhibitor, UO126, markedly reduces lipid deposition in RPMs treated with 50 mg/l Ox-LDL alone or in combination with 10 μ M UO126 for 12, 24 and 48 h. (A) Cells were stained with oil red O and examined under an inverted microscope (magnification, x400). RPMs were observed to engulf a large quantity of lipids (red). (B) No red fluorescence was detected in untreated RPMs, whereas red fluorescence was detected in DiI-Ox-LDL-treated RPMs, suggesting lipid deposition in macrophages (magnification, x400). ERK, extracellular signal-regulated kinases; RPMs, rat peritoneal macrophages; Ox-LDL, oxidized-low-density lipoprotein.

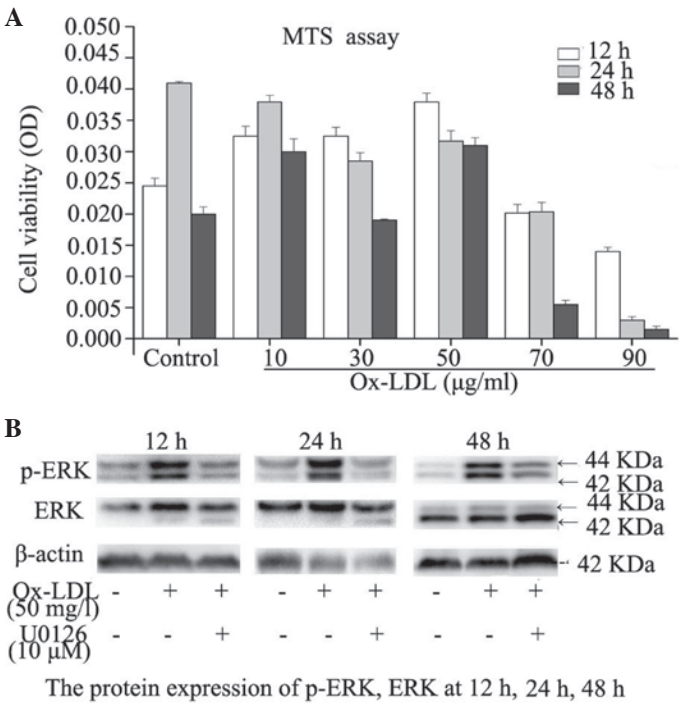


Figure 2. Effects of Ox-LDL treatment on ERK and p-ERK protein expression levels in RPMs. (A) RPMs were treated with Ox-LDL at various concentrations for 12, 24 and 48 h. Cell viability was determined by MTS assay. OD demonstrates the viability of cells. (B) RPMs were treated with 50 mg/l Ox-LDL alone or in combination with 10 μ M UO126 (ERK inhibitor) for 12, 24 and 48 h, and western blotting was performed to detect ERK and p-ERK protein expression levels. Data are presented as the mean \pm standard deviation. OD, optical density; Ox-LDL, oxidized-low-density lipoprotein; RPMs, rat peritoneal macrophages; ERK, extracellular signal-regulated kinases; p-ERK, phosphorylated ERK.

no significant difference was detected in ABCG1 mRNA and protein expression levels at 48 h post-treatment with Ox-LDL alone or in combination with UO126, and the mRNA and protein expression levels of ABCG1 were not higher in RPMs treated with Ox-LDL + UO126, as compared with RPMs treated with Ox-LDL alone. These results indicate that

ERK1/2 inhibition augments mRNA and protein expression levels of ABCG1 in Ox-LDL-stimulated macrophages.

Inhibition of ERK1/2 reduces CD36 expression in Ox-LDL-treated macrophages. The effect of ERK1/2 inhibition on Ox-LDL-induced CD36 expression was evaluated by

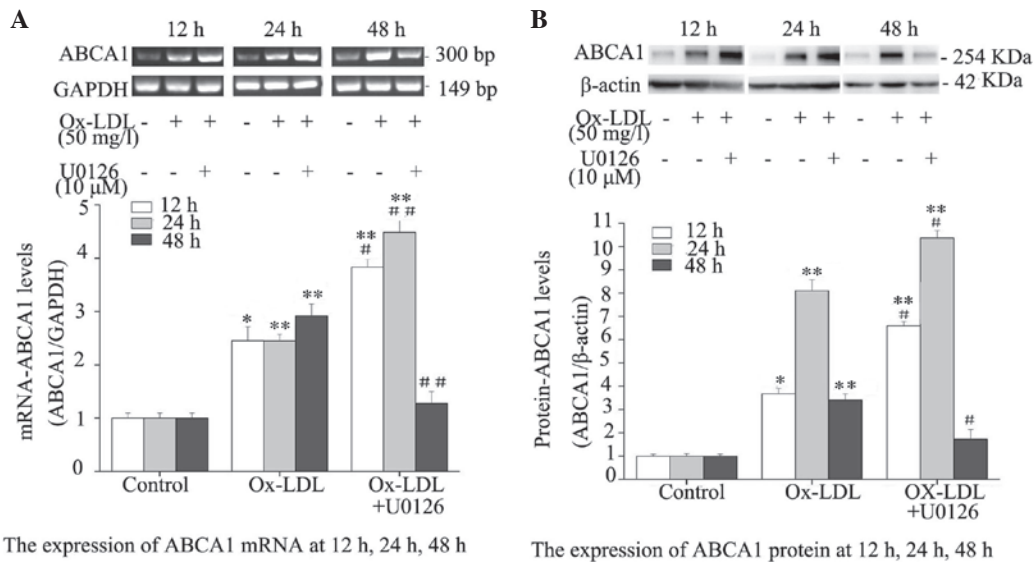


Figure 3. ABCA1 mRNA and protein expression levels in RPMs. RPMs were treated with 50 mg/l Ox-LDL alone or in combination with 10 μM U0126 (ERK inhibitor) for 12, 24 and 48 h. (A) Reverse transcription-polymerase chain reaction was used to determine ABCA1 mRNA expression. GAPDH served as an internal control. (B) Whole cell lysates were used for western blot analysis to determine ABCA1 protein expression. Data are presented as the mean ± standard deviation. *P<0.05, **P<0.01, the Ox-LDL group vs. the control group; #P<0.05, ##P<0.01, the Ox-LDL + U0126 group vs. the Ox-LDL group. Ox-LDL, oxidized-low-density lipoprotein; ABCA1, ATP-binding cassette transporter A-1; RPMs, rat peritoneal macrophages; GAPDH, glyceraldehyde 3-phosphate dehydrogenase.

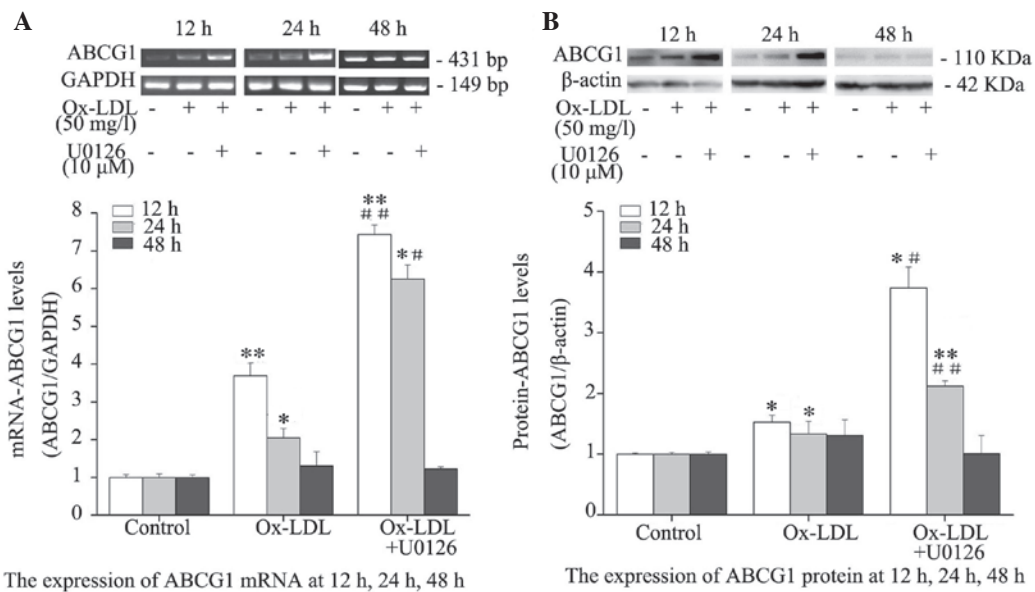


Figure 4. Effects of ERK1/2 inhibitor, U0126, on ABCG1 mRNA and protein expression levels in RPMs. RPMs were treated with 50 mg/l Ox-LDL alone or in combination with 10 μM U0126 for 12, 24 and 48 h. (A) Reverse transcription-polymerase chain reaction was used to determine ABCG1 mRNA expression. GAPDH served as an internal control. (B) Whole cell lysates were used for western blot analysis to determine protein expression. Data are presented as the mean ± standard deviation. *P<0.05, **P<0.01, the Ox-LDL group vs. the control group; #P<0.05, ##P<0.01, the Ox-LDL + U0126 group vs. the Ox-LDL group. RPMs, rat peritoneal macrophages; ERK, extracellular signal-regulated kinases; Ox-LDL, oxidized-low-density lipoprotein; ABCG1, ATP-binding cassette transporter G-1; GAPDH, glyceraldehyde 3-phosphate dehydrogenase.

RT-PCR and western blotting. Treatment with Ox-LDL resulted in a significant elevation in the expression levels of CD36, as compared with the untreated controls (P<0.05), and co-treatment with Ox-LDL and U0126 for 12, 24 and 48 h resulted in significant reductions in mRNA (Fig. 5A) and protein (Fig. 5B) expression levels of CD36, as compared with treatment with Ox-LDL alone. These results suggest that Ox-LDL markedly induces CD36 expression and inhibition of ERK1/2 reduces CD36 expression in Ox-LDL-treated macrophages.

Discussion

Atherosclerotic lesions are characterized by accumulation of cholesterol in the arterial tunica intima. Macrophages are central to the development and progression of AS (13,14); they have been observed to remove cholesterol deposits in arteries, which is beneficial in the initial stages of AS; however, continued cholesterol accumulation may result in the formation of foam cells, which are involved in AS progression (15,16).

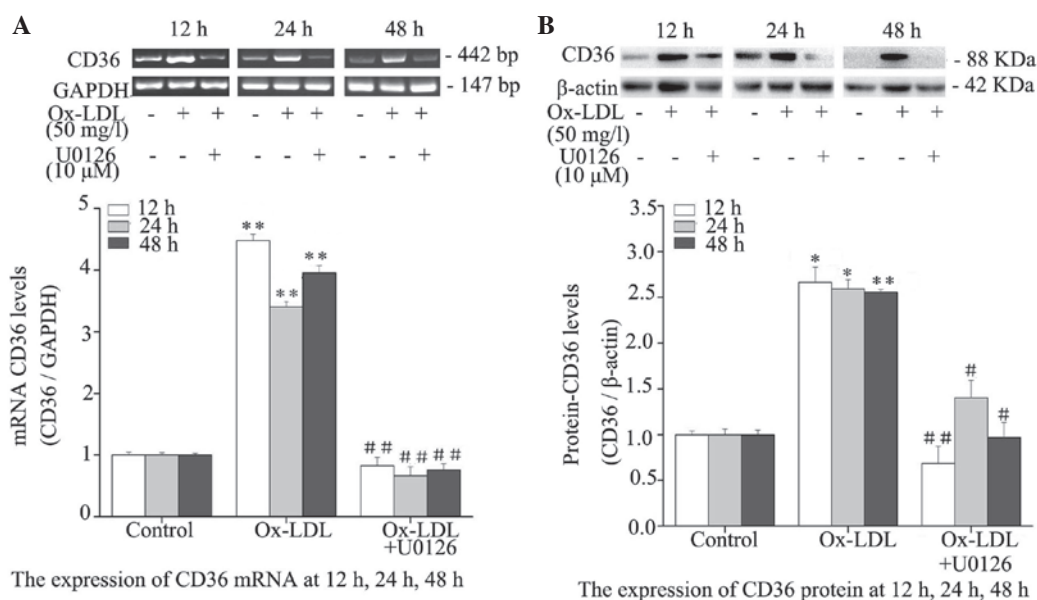


Figure 5. Effects of ERK1/2 inhibitor, U0126, on CD36 mRNA and protein expression levels in RPMs. RPMs were treated with 50 mg/l Ox-LDL alone or in combination with 10 μ M U0126 for 12, 24 and 48 h. (A) Reverse transcription-polymerase chain reaction was used to determine CD36 mRNA expression. GAPDH served as an internal control. (B) Whole cell lysates were used for western blot analysis to determine CD36 protein expression. Data are presented as the mean \pm standard deviation. * P <0.05, ** P <0.01, the Ox-LDL group vs. the control group; # P <0.05, ## P <0.01, the Ox-LDL + U0126 group vs. the Ox-LDL group. CD, cluster of differentiation; Ox-LDL, oxidized-low-density lipoprotein; RPMs, rat peritoneal macrophages; ERK, extracellular signal-regulated kinases; GAPDH, glyceraldehyde 3-phosphate dehydrogenase.

The response of RPMs to modified LDL is considered a good model for AS research (17,18). The results of the present study indicated that RPMs engulfed a large quantity of Ox-LDL, as demonstrated by DiI fluorescence and oil red O staining. In addition, the present study demonstrated that inhibition of ERK1/2 markedly reduced Ox-LDL deposition in the RPMs. It has previously been reported that ERK1/2 is associated with cell migration, survival, differentiation and proliferation (4). However, the association between ERK1/2 and lipid metabolism remains to be elucidated.

Cholesterol uptake is a pathway by which extracellularly modified LDLs are ingested by macrophages via scavenger receptors. As a major scavenger receptor of Ox-LDL (19,20), CD36 binds to >50% of Ox-LDL in macrophages (21), which is crucial to the progression of AS. Absence of CD36 on the cell surface of macrophages has been demonstrated to be protective against AS (22). In addition, treatment with a CD36 competitive peptide ligand (EP80317) that blocks the Ox-LDL-binding site of CD36 resulted in a marked reduction (up to 51%) of atherosclerotic lesions in apolipoprotein E-deficient mice (23). It has therefore been hypothesized that inhibition of CD36 expression in macrophages has potential anti-atherosclerotic effects. It has been reported that the CD36 signaling pathway may be initiated by internalization of Ox-LDL (3,7). Consistent with these previous findings, the present study indicated that Ox-LDL resulted in a notable increase in CD36 expression. Increasing evidence suggests that Ox-LDL increases ERK phosphorylation (3,20,24). The results of the present study indicated that inhibition of ERK1/2 markedly reduced Ox-LDL-induced CD36 expression in RPMs. Furthermore, a p38 MAPK inhibitor, SB203580, has previously been demonstrated to suppress CD36 expression in THP-1 cell lines (24) and RAW264.7 cells (20). The present study demonstrated

that treatment with an ERK1/2 inhibitor, U0126, attenuated Ox-LDL-induced CD36 expression at the transcriptional and translational level. It is therefore assumed that ERK1/2 may be important in mediating CD36 expression in macrophages.

Macrophages have been reported to maintain cellular lipid homeostasis via cholesterol efflux pathways. The principle molecules associated with cholesterol efflux in macrophages are ABCA1 and ABCG1 (25,26). In THP-1 cells, Ox-LDL has been shown to upregulate ABCA1 expression (27). The present study demonstrated that the mRNA and protein expression levels of ABCA1 were significantly increased following Ox-LDL treatment, and that inhibition of ERK1/2 increased ABCA1 mRNA and protein expression levels in macrophages. It has been reported that ERK1/2 inhibitors synergize with LXR activation, in order to induce ABCA1 expression in a RAW macrophage cell line (5), and U0126, an ERK1/2 inhibitor, has been demonstrated to delay the degradation of ABCA1 mRNA and protein in macrophages (28). However, a previous study indicated that inhibition of ERK1/2 was able to promote ABCA1 and ABCG1 protein degradation in tumor cells, whereas inhibition of ERK1/2 upregulated ABCA1 and ABCG1 expression in human THP-1 macrophages (29). It is therefore hypothesized that the effect of ERK1/2 inhibition on ABCA1 and ABCG1 expression may depend on cell specificity. The results of the present study demonstrated that inhibition of ERK1/2 increased ABCA1 expression in Ox-LDL-induced macrophages at 12 and 24 h, and gradually decreased ABCA1 expression at 48 h. The half-life of ABCA1 protein is 1-2 h, and ABCA1 is frequently present during homeostasis between expression and degradation. Inhibition of ERK1/2 has been demonstrated to suppress the degradation of ABCA1 mRNA and protein (5) or increase its stability (5). However, macro-

phages secrete various inflammatory cytokines, which affect ABCA1 expression (30,31). Further studies are required to investigate the mechanisms underlying the effects of ERK1/2 inhibition on the regulation of cytokines that affect ABCA1 expression. The present study demonstrated that alterations to ABCG1 mRNA and protein expression levels were similar to those observed in ABCA1 expression at 12 and 24 h post-treatment with Ox-LDL. It has been reported that macrophages may overexpress unsaturated fatty acids and products of 12/15-lipoxygenase, which increase the degradation of ABCG1 protein (32,33). In addition, ABCG1 expression is affected by cell activity and cytokines secreted from macrophages (34).

The present study indicated that inhibition of ERK1/2 markedly suppressed lipid deposition in macrophages by promoting ABCA1 and ABCG1 expression, which are associated with cholesterol efflux. ABCA1 and ABCG1 expression may be induced by ERK1/2 inhibition or LXR α (35), thus indicating that an association may exist between LXR and ERK. Since ABCA1 and ABCG1 are nuclear receptor LXR target genes involved in cholesterol efflux, activation of LXR may increase ABCA1 and ABCG1 expression in macrophages (36,37). However, it has been reported that ERK1/2 inhibitors have no effect on LXR α and LXR β expression (5). It is conceivable that ERK1/2 inhibition may directly increase ABCA1 and ABCG1 expression by methods other than upregulating LXR expression. ERK1/2 controls transcriptional and post-transcriptional regulation of LXR and peroxisome proliferator-activated receptor (PPAR) α/γ (29,38-40), and mediates expression of their target genes. Further studies are required to elucidate the association between ERK and PPARs/LXRs-ABCA1/G1, and to understand the underlying mechanisms.

In conclusion, inhibition of ERK1/2 increases macrophage lipid efflux by upregulating ABCA1 and ABCG1 expression, and suppresses lipid engulfment by downregulating CD36 expression at the transcriptional and translational level. These findings suggest that inhibition of ERK1/2 may exert potential anti-atherosclerotic effects, and involvement of the ERK1/2 pathway in lipid metabolism may provide additional knowledge for the development of novel treatment strategies for AS.

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