Dilazep, an Antiplatelet Agent, Inhibits Tissue Factor Expression in Endothelial Cells and Monocytes

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Dilazep, an antiplatelet agent, is generally used as an anti-thrombotic drug in clinical practice. Dilazep is also known to exert cytoprotective and antioxidant effects on endothelial cells. However, its effect on the endothelial or monocyte procoagulant activity is unknown. In the current study, the effect of dilazep on the expression of tissue factor (TF) in human umbilical vein endothelial cells (HUVECs) after the stimulation with tumor necrosis factor-α (TNF), thrombin, or phorbol 12-myristate 13-acetate (PMA) was evaluated. We also evaluated the effect of dilazep on TNF (1,000 U/mL)-induced TF expression on monocytes. Dilazep inhibited TF activity induced on HUVECs by each stimulant, TNF (1,000 U/mL), thrombin (25 nmol/L), or PMA (5 nmol/L) in a dose-dependent fashion (1 to 100 µg/mL). TF activity decreased to approximately 10% after treating with 100 µg/mL of dilazep. Dilazep also blocked the expression of TF antigen induced by each stimulant on the surface of HUVECs as determined by flow cytometric analysis. In addition, in HUVECs, it significantly decreased the expression of TF mRNA and the total TF antigen induced by thrombin or PMA, but not those induced by TNF, suggesting that dilazep blocks the TF expression induced by PMA or thrombin at a transcriptional and posttranscriptional level and that induced by TNF at a posttranscriptional level. Western blot analysis showed that dilazep reduces the accumulation of native TF but increases that in lower molecular weight TF derivatives. The adenosine receptor antagonist, 8-(p-sulphonyl)theophylline, partially counteracted the anticoagulant activity of dilazep on HUVECs, thereby suggesting that the inhibitory effect of dilazep on TF expression in HUVECs depends, at least in part, on its adenosine potentiating activity. Dilazep also inhibited TNF-induced TF expression on monocytes in a dose-dependent fashion (0.1 to 100 µg/mL). In brief, the current study showed for the first time that dilazep, a commonly used antiplatelet drug, strongly inhibits the TF expression in HUVECs and monocytes. Dilazep may have a potent therapeutic value in patients with hypercoagulable state for its inhibitory property on the procoagulant activity of endothelial cells and monocytes.

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MATERIALS AND METHODS

Materials. Citrated human plasma was obtained from the local Red Cross blood center. Human factor VIIa and factor X were provided by Dr Tomohiro Nakagaki (Chemo-Sera Therapeutic Institute, Kumamoto, Japan). Antithrombin, protein C, and thrombin were prepared from human plasma as described. Recombinant human TF was isolated from the Asahi Chemical Industry (Fujii, Shizuoka, Japan). The synthetic peptide substrate for factor Xa, N-benzyol-L-isoleucyl-L-glutamyl-glycyl-L-arginine-p-nitroanilide hydrochloride (S2222), was purchased from Chromogenix (Malmö, Sweden). The fluorogenic substrate for activated protein C, Boc-Leu-Ser-Thr-Arg-methylcoumarylaldehyde (MCA), was obtained from the Protein Research Foundation (Osaka, Japan). Dulbecco’s phosphate-buffered saline (PBS), gentamycin, bovine serum albumin (BSA), heparin (174 U/mg), and adenosine were from Sigma (St

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Louis, MO). Nick column was obtained from Pharmacia-LKB (Uppsala, Sweden). Triton X-100 and aprotonin were from Boehringer Mannheim GmbH (Mannheim, Germany). Glyceraldehyde-3 phosphate dehydrogenase (GAPDH) cDNA probe and 8-(p-sulfophenyl) theophylline (8-SPT) was purchased from Clontech Labs (Palo Alto, CA) and Research Biochemicals International (Natick, MA), respectively. Horseradish peroxidase (HRP)-conjugated antimonute IgG-goat antibody and antigen IgG-rabbit antibody were obtained from Bio-Rad (Richmond, CA). The reagents to check the viability of the cells, WST-1 [2-(4-iodophenol)-3-(4-nitrophenol)-5-(2,4-disulfo- phenyl)-2H-tetrazolium-Na] and 1-methoxy PMS (1-methoxy-5-methyl-phenazine methylsulphate), were purchased from Dojindo (Kumamoto, Japan).

Patients. Twenty inoperable patients with malignant disease (adenocarcinoma, 12; squamous cell carcinoma, 2; small-cell carcinoma, 2; hepatoma, 2; melanoma, 1; and myeloma, 1) and 8 with benign disorder (dissecting aneurysm of the aorta, 6; chronic enteritis, 1; and chronic viral infection, 1) were studied. All of these patients were in the hypercoagulable state and were treated with dila-zep (300 mg/d) for 1 month before the initiation of any kind of therapy. Blood samples were collected from the patients before and after 1 month of starting the therapy with dila-zep. Plasma levels of thrombin-antithrombin complex (TAT) and fibrin degradation product (FDP)-D-dimer were determined using Enzygnost-TAT kit (Behringwerke AG, Marburg, Germany; normal range, 1.8 ± 0.7 ng/mL) and Felsia D-dimer (Agen, Brisbane, Australia; normal range, 48 ± 34 ng/mL), respectively. Plasma fibrinogen level was determined using the clotting time method (normal range, 2.5 ± 0.4 mg/mL).

The plasma level of TF antigen was determined by TF enzyme-linked immunosorbent assay (ELISA) kit (Chemo-Sera Therapeutic Institute; normal range, 196.8 ± 54.3 pg/mL). Data are expressed as the mean ± SD. The statistical difference between variables was evaluated by the Student's t-test or the Wilcoxon rank test according to the Gaussian distribution of the data.

Culture of HUVECs. HUVECs were purchased from Kurabo (Osaka, Japan). The cells were cultured in modified MCD medium (Chlorella Industry, Tokyo, Japan) supplemented with 2% fetal bovine serum (GIBCO BRL, Grand Island, NY), 50 μg/mL gentamycin, 0.01 μg/mL epithelial growth factor (Becton Dickinson, Bedford, MA), 10 μg/mL endothelial cell growth factor (Becton Dickinson), and 10 μg/mL unfracthioned heparin (Sigma) under an atmosphere of 95% air and 5% CO2.

Preparation of peripheral blood mononuclear cells. Peripheral blood cells were obtained from healthy donors by vein puncture using EDTA as anticoagulant. Mononuclear cells were isolated by the Lymphoprep Tube (Nycomed Pharma Diagnostica, Oslo, Norway). The mononuclear cell phase, comprising monocytes and lymphocytes, was harvested, washed twice with RPMI (RPMI 1640 medium supplemented with 2 mM L-glutamine, 100 μg/mL streptomycin, 100 μg/mL penicillin, 10% fetal bovine serum), and resuspended in RPMI.

Determination of HUVEC surface TF activity. TF activity was determined as factor X activation by factor VIIa/TFA complex on HUVECs after stimulation with TNF, thrombin, or PMA. After incubation of the cells (24-well plate, 5 × 105 cells/well) with each stimulant in the presence or absence of dilazep or adenosine, the cells were washed twice with HEPES-buffered saline (20 mM/L HEPES, 150 mM/L NaCl, pH 7.5) containing 5 mM/L CaCl2 and 10% fetal bovine serum, and resuspended in RPMI.

Analysis of TF mRNA with reverse transcription and polymerase chain reaction (RT-PCR). Total RNA of HUVECs was prepared using the RNA zol (Cinna/Biotex Laboratories, Houston, TX) after stimulating the cells with TNF, thrombin, or PMA in the presence or absence of dilazep. RT-PCR was performed as described previously.

Flow cytometric analysis of HUVEC surface TF antigen. HUVECs were incubated with or without TNF, thrombin, or PMA in the presence or absence of dilazep for 1 hour. After washing the cells twice with cold Dulbecco’s PBS and resuspended (2 × 105 cells) in 100 μL PBS, the cells were incubated with mouse monoclonal anti-human TF IgG (5 μg/mL) for 30 minutes at 4°C and subsequently washed three times with PBS. The cells were then incubated with the F(ab')2 fraction of a fluorescein isothiocyanate (FITC)-labeled antimonute IgG-goat antibody (1/20 dilution) for 30 minutes at 4°C. The cells were then washed three times with PBS and fixed in fluorescence-activated cell sorting (FACS) lysing solution (Becton Dickinson) according to the manufacturer’s instructions. Flow cytometric analysis was performed using a FACScan analyzer (Becton Dickinson) equipped with a 15-mW, 488-nm argon-ion laser. Green fluorescence emission was recorded at 530 nm. A total of 10,000 events were analyzed per sample. Median channel fluorescence in each sample was determined using a logarithmic scale. Logarithmic channel was converted into linear channel fluorescence units to facilitate comparison of results.

Measurement of total TF antigen level in HUVECs. The total TF antigen levels in lysates of HUVECs were measured by ELISA. Briefly, after stimulation of HUVECs with TNF, thrombin, or PMA in the presence or absence of dilazep, the cells were washed twice with cold Dulbecco’s PBS and harvested. Subsequently, harvested cells were centrifuged (2,000 rpm for 5 minutes) and the precipitate was dissolved in Tris-buffered saline (TBS; 50 mM/L Tris/HC1, 150 mM/L NaCl, pH 7.5) containing, 0.6% (wt/vol) Triton X-100 and aprotonin (100 μ/mL). The cellular extraction was then diluted 1,000-fold with TBS containing 0.1% BSA and 0.02% thimerosal and applied to wells coated with rabbit antihuman TF antibodies (American Diagnostica, Greenwich, CT). The wells were blocked with 100 mM/L phosphate buffer, pH 8.0, containing 5% BSA, and then washed three times with TBS containing 0.1% BSA and 0.02% thimerosal. Then, biotynyl-monoclonal antibody against human TF (MSTF-3) was added and incubated for 2 hours. After washing the wells, HRP-labeled streptavidin was added and incubated for 1 hour. After the addition of H2O2 and peroxidase substrate, color development was measured at 492 nm using an EAR microplate reader.

Western blotting analysis of TF antigen in HUVECs. TF expression in HUVECs was determined by Western blotting using polyclonal goat antihuman TF antibodies (American Diagnostica). Sodium dodecyl sulphate-polyacrylamide gel electrophoresis of HUVECs lysates was performed, followed by blotting onto nitrocellulose membrane. The membrane was incubated with blocking solution (100 mM/L PBS, pH 7.5, containing 5% BSA and 0.5% Tween 20) and then exposed to a 1/2,000 dilution of goat anti-TF antibodies for 1 hour while shaking at room temperature. After washing the membrane with PBS containing 0.1% BSA and 0.5% Tween 20, the membrane was then incubated with a 1/20,000 dilution of antigoat IgG antibody conjugated to HRP. The membrane was washed and quantitated using an enhanced chemiluminescence detection system (ECL; Amersham, Buckinghamshire, UK) as described previously.

Analysis of TF mRNA with reverse transcription and polymerase chain reaction (RT-PCR). Total RNA of HUVECs was prepared using the RNA zol (Cinna/Biotex Laboratories, Houston, TX) after stimulating the cells with TNF, thrombin, or PMA in the presence or absence of dilazep. RT-PCR was performed as described previously. Briefly, total RNA (1 μg) was reverse transcribed by superscript preamplification system (GIBCO BRL) with random primes and amplified using PCR.
hexamer primers, and the reverse-transcribed cDNA was then amplified by PCR using the primers based on the mRNA of TF or GAPDH, a housekeeping gene that was not affected by the stimuli used in this study. The sequences of the primers used for TF mRNA amplification were 5’ ACTACTGTATCTGGTCTCCTCCTAGCTGACCGTTC-3’ corresponding to 722-751 nucleotides and the 5’ ATTCACTGGGGATTCCTCCTGAGTGGCACC 3’ corresponding to 925-954 nucleotides. The sequences of the primers used for GAPDH mRNA amplification were 5’ CCACCCATGCAAAATTCCATGCA-3’ corresponding to 150-169 nucleotides and 5’ TCTAGACGGCAGGTCAGGTCCACC 3’ corresponding to 720-743 nucleotides. PCR was performed with 22 cycles of 1 minute of denaturation at 94°C, 1 minute of hybridization at 52°C, and 2 minutes of elongation at 73°C. An aliquot of sample was loaded onto a 2% agarose gel. Amplified DNA of TF or GAPDH was blotted on a nylon membrane and hybridized with the TF or GAPDH cDNA probe labeled with a [32P]dCTP (3,000 Ci/mmol) using a DNA labeling kit (TaKaRa Shuzo, Osaka, Japan). The membrane was then exposed to x-ray film at −70°C. The intensity of the hybridization signals was densitometrically quantified by a Fujix BAS-2000 Bio-Image analyzer (Fuji Photo Film, Tokyo, Japan). The amount of TF mRNA was normalized against the GAPDH mRNA.

**Determination of HUVEC surface thrombomodulin (TM) activity and TM antigen in HUVECs.** TM activity on the surface of HUVECs was determined as a cofactor activity of TM for thrombin-catalyzed protein C activation using 48-well collagen-coated plates, as described previously. After washing the HUVECs (2 × 10⁶ cells) in the wells three times, 50 μL of TBS, pH 8.0, containing 0.1% BSA and 5 mmol/L CaCl₂, 25 μL of 0.5 μg/mL thrombin, and 50 μL of 50 μg/mL protein C was added to each well. After 1 hour of incubation at 37°C, antithrombin (5 μg/mL, final concentration) and heparin (2 U/mL, final concentration) were added to the wells. The amount of activated protein C in the reaction mixture (100 μL) was assayed by incubating with 2 mL of 200 μmol/L Boc-Leu-Ser-Thr-Arg-MCA in TBS. The fluorescence of the released amino-methylcoumarin was determined using a fluorescence spectrophotometer (Shimadzu, Kyoto, Japan) with excitation at 380 nm and emission at 440 nm.

The amount of total TM antigen in HUVECs was determined by an ELISA using two murine monoclonal antihuman TM IgGs as reported previously. Briefly, HUVECs were incubated with 100 μg/mL dilazep, followed by appropriate washing with cold Dulbecco’s PBS and harvesting of the cells. Subsequently, the cells were centrifuged (2,000 rpm for 5 minutes) and the precipitate was dissolved in 50 μL TBS containing 0.6% (wt/vol) Triton X-100 and 0.2 U/mL aprotinin. The cell extract was then applied to the wells coated with a monoclonal anti-TM IgG (MFTM-5). The wells were blocked with PBS, pH 8.0, containing 5% BSA and then incubated for 5 minutes with BSA containing 0.1% BSA and 0.02% thimerosal. Thereafter, HRP-labeled monoclonal anti-TM IgG (MFTM-6) was added and incubated for 1 hour. After the addition of H₂O₂ and peroxidase substrate, color development was measured at 492 nm using an EAR microplate reader.

**Determination of monocytic cell surface TF activity.** Mononuclear cells (monocytes plus lymphocytes) suspended in RPMI were incubated with or without 100 μg/mL dilazep for 15 minutes and stimulated with TNF for 5 hours. Procoagulant activity (PCA) on the surface of intact monocytes was determined by a single-stage clotting assay as described previously. Briefly, 50 μL of mononuclear cell suspension was mixed with 50 μL of 25 mmol/L CaCl₂, Clotting was initiated by the addition of 50 μL of normal human plasma, and the clotting time at 37°C was recorded using KC 10 coagulometer (Heinrich Amelung, GmbH, Lemgo, Germany). The PCA was extrapolated from a standard curve drawn using rabbit brain thromboplastin standard (Ortho, Raritan, NJ), as described previously. Cell surface-associated PCA was linear over the range of 12 × 10⁵ mononuclear cells/mL. PCA of monocytes was confirmed to depend on TF, because it was specifically inhibited by polyclonal anti-TF antibody.

**RESULTS**

**Effect of dilazep in patients with hypercoagulable state.** We evaluated the effect of dilazep on hemostatic markers in patients with hypercoagulable state. The plasma levels of TAT (32.3 ± 19.7 μg/mL), FDP-D-dimer (1,411.1 ± 384.4 μg/mL), fibrinogen (4.1 ± 0.1 mg/mL), and TF antigen (355.5 ± 94.0 μg/mL) significantly (P < .001) decreased after treatment with dilazep (300 mg/d), as shown in Fig 1.
Fig 2. Time course of TF activity expression on HUVECs stimulated with TNF, thrombin, or PMA and the effect of dilazep on the TF expression. HUVECs (5 × 10^5/well) were stimulated with (A) 1,000 U/mL TNF, (B) 25 nmol/L thrombin, or (C) 5 nmol/L PMA in the presence (●) or absence (○) of 100 μg/mL dilazep. The TF activity on HUVEC surface was determined at 0, 3, 5, 6, 9, 12, and 24 hours after incubation with each stimulant as factor Xa generation (A405nm /20 minutes) using S2222 (see the Materials and Methods). The values represent the mean ± SD of four independent experiments.

These hemostatic parameters also decreased significantly in each patient group (malignant and benign disease) after treatment with dilazep (P < .001) as compared with baseline values (data not shown). This finding suggests that dilazep has a potent anticoagulant activity on endothelial cells or intravascular leukocytes. This preliminary clinical study prompted us to evaluate the effect of dilazep on endothelial cell procoagulant activity.

Effect of dilazep on TF activity of HUVECs stimulated with TNF, thrombin, or PMA. We evaluated the effect of dilazep on TF expression in HUVECs stimulated with the inflammatory mediators TNF, thrombin, or PMA. A gradual increase in TF activity was observed on HUVECs after stimulating with TNF, thrombin, or PMA, reaching a maximum level after 5 to 6 hours. As shown in Fig 2, 100 μg/mL dilazep inhibited this stimulatory effect of TNF, thrombin, or PMA on TF activity of HUVECs. TF activity on stimulated HUVECs for 5 hours depended on the dose of the stimulant as described previously, and the activity reached maximal levels at concentrations of 1,000 U/mL TNF, 25 nmol/L thrombin, and 5 nmol/L PMA, respectively. Coincubation with 100 μg/mL dilazep decreased approximately 90% of the TF activity induced by each stimulant (Fig 3). Increasing concentrations of dilazep (0 to 100 μg/mL) decreased the cell surface TF activity in a dose-dependent fashion (Fig 4). The minimal concentration of dilazep required for inhibiting the TNF-stimulated TF expression was 1 μg/mL, but that for the thrombin- and PMA-stimulated TF expression was 0.1 μg/mL. The concentrations of dilazep required for 50% inhibition of the expression of TF activity on HUVECs induced by 1,000 U/mL TNF, 25 nmol/L thrombin, and 5 nmol/L PMA were approximately 20, 2.5, and 20 μg/mL.

Fig 3. The effect of dilazep on TF activity on HUVECs stimulated with varying concentrations of TNF, thrombin, or PMA. HUVECs were treated for 5 hours with varying concentrations of each stimulant. (A) TNF (0 to 2,500 U/mL), (B) thrombin (0 to 250 U/mL), and (C) PMA (0 to 250 nmol/L) in the presence (●) or absence (○) of 100 μg/mL dilazep. TF activity on HUVEC surface was determined as factor Xa generation (A405nm /20 minutes) using S2222 (see the Materials and Methods). The values represent the mean ± SD of four independent experiments.
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respectively. The viability of HUVECs as determined by using WST-1 and 1-methoxy PMS was not affected by dilazep at any concentration used in this experiment.

To discern whether the inhibitory effect of dilazep on TF expression is caused by a global effect on protein synthesis, the effect of dilazep on TM, which is constitutively expressed on resting HUVECs, was also investigated. The time course of the levels of TM activity and the total TM antigen expressed on HUVECs in the presence (100 μg/mL) or absence of dilazep were determined. The TM activity and the total TM antigen levels in the cells were not affected by dilazep even after 24 hours of the treatment (data not shown).

**Flow cytometry of TF antigen expression.** FACScan analysis was performed to determine whether dilazep reduces the amount of TF antigen expression on HUVEC surface induced by TNF, thrombin, or PMA. As shown in Fig 5, dilazep alone did not affect the fluorescence intensity on HUVECs, but markedly decreased the elevation of TF antigen levels on HUVECs induced by 1,000 U/mL TNF, 25 nmol/L thrombin, or 5 nmol/L PMA.

**Effect of dilazep on total TF antigen level in HUVECs stimulated with TNF, thrombin, or PMA.** To investigate whether the suppressive effect of dilazep on HUVEC surface TF activity depends on the decreased TF production by the cells, the total amount of TF antigen in HUVEC lysates was measured by ELISA. Uprogelation of total TF antigen level was observed in HUVECs treated with 1,000 U/mL TNF, 25 nmol/L thrombin, and 5 nmol/L PMA. Dilazep (100 μg/mL) significantly inhibited the thrombin- and PMA-induced upregulation of total TF antigen levels (432% ± 31.4% vs 158% ± 0.8% and 526% ± 12.9% vs 210% ± 4.6%, respectively). However, dilazep did not inhibit the TNF-induced total TF antigen level (425% ± 1.1% vs 405% ± 14.2%; Fig 6). Considering this latter result together with that shown in Fig 5B, dilazep appears to inhibit the transport of TFN-induced TF protein to the cell surface of HUVECs.

**Effect of dilazep on the molecular mass of TNF-induced TF in HUVECs.** To investigate the inhibitory effect of dilazep on TNF-induced TF expression on HUVECs, Western blot analysis was also performed. TNF-stimulated HUVECs exhibited a prominent band with approximate molecular mass of 47 kD, consistent with an increase in the mature form of TF. There was also an increase in a band at the 33-kD level. The addition of dilazep to TNF stimulation prevented the increase in the high molecular mass band (47 kD) but caused an increase in the band at the 33-kD level (Fig 7).

**Effect of dilazep on TF mRNA expression induced by TNF, thrombin, or PMA in HUVECs.** To analyze whether the effect of dilazep on TF antigen levels in HUVECs (stimulated with TNF, thrombin, or PMA) corresponds to the levels of TF mRNA in HUVECs, the effect of dilazep on the steady-state levels of TF mRNA was assessed in HUVECs using the RT-PCR method.27 Resting HUVECs and the cells treated with dilazep alone showed no detectable expression of TF mRNA (Fig 8A). As shown in Fig 8A and B, 1,000 U/mL TNF, 25 nmol/L thrombin, or 5 nmol/L PMA induced TF mRNA expression markedly after 3 hours of the treatment with each stimulant. Dilazep significantly inhibited the thrombin- or PMA-induced TF mRNA expression. However, the same amount of dilazep did not decrease the TF mRNA expression induced by TNF. These effects of dilazep on the TF mRNA expression appear to correspond well to that on the total TF antigen level in HUVECs shown in Fig 6.

**Effect of adenosine receptor antagonist on the inhibitory effect of dilazep.** Dilazep is believed to potentiate the effect of adenosine by blocking the uptake of adenosine.31 In an attempt to clarify whether the inhibitory effect of dilazep on TF expression is mediated by adenosine receptor, the effect of the adenosine receptor antagonist, 8-SPT, on the inhibitory effect of dilazep was investigated. As shown in Fig 9, 1 mmol/L 8-SPT alone did not affect the basal or stimulant-induced TF expression. Dilazep (100, 10, and 100 μg/mL) decreased the TF activity induced by 1,000 U/mL TNF, 25 nmol/L thrombin, and 5 nmol/L PMA, respectively. Down-regulation of TF activity on HUVECs induced by dilazep was counteracted by 8-SPT, indicating that the inhibitory effect of dilazep on TF expression on HUVECs is mediated, at least in part, by adenosine receptor.
Fig 5. Flow cytometry of the effect of dilazep on TF antigen expression on the surface of HUVECs stimulated with TNF, thrombin, or PMA. HUVECs were stimulated for 5 hours with each stimulant. (A) Control, (B) 1,000 U/mL TNF, (C) 25 nmol/L thrombin, and (D) 5 nmol/L PMA, in the presence (p) or absence (a) of 100 μg/mL dilazep. The cells were then harvested, incubated with murine monoclonal antihuman TF IgG, and then incubated with FITC-labeled goat antimouse IgG-antibody. Fluorescence intensity was determined by FACSscan as described in the Materials and Methods, and the data are displayed using a logarithmic scale.

Effect of adenosine on the TF activity of HUVECs stimulated with TNF, thrombin, or PMA. Increasing concentrations of adenosine (0 to 10 μmol/L) decreased the TF activity expression on the surface of HUVECs stimulated with 1,000 U/mL TNF, 25 nmol/L thrombin, or 5 nmol/L PMA in a dose-dependent fashion (Fig 10). Coincubation with 10 μmol/L adenosine decreased approximately 50% the TF activity on HUVECs induced by each stimulant. These findings indicate that adenosine significantly blocks the TF expression in HUVECs stimulated with TNF, thrombin, or PMA. Viability of HUVECs as determined using WST-1 was not affected by adenosine at any concentration used in this experiment.

Additive effect of dilazep and adenosine on the TF activity of HUVECs stimulated with TNF, thrombin, or PMA. Isotope-technical studies using ³H-dilazep in humans showed that the therapeutic concentration of dilazep in the systemic circulation is around 1 μg/mL. The therapeutical concentration of dilazep is known to increase the concentration of adenosine in the extracellular space fluid up to micromolar levels. In our assays, we used a dose of dilazep equal to its therapeutic blood concentration in combination with adenosine. We found that dilazep, in combination with adenosine, inhibited the 1,000 U/mL TNF- and 25 nmol/L thrombin-induced TF expression in HUVECs more significantly (60%) than dilazep alone, as shown in Fig 11.

Effect of dilazep on TF activity of monocytes stimulated with TNF. Peripheral blood monocytes are known to participate in blood coagulation processes by expressing TF after being stimulated with inflammatory mediators. The
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Viability of mononuclear cells as determined by using WST-1 and 1-methoxy PMS was not affected by dilazep at any concentration used in this experiment.

DISCUSSION

Various cytokines, such as TNF, IL-1, and lymphotokin, play relevant roles in several inflammatory processes and in the pathogenesis of atherosclerosis. Activation of endothelial cells by these mediators induces the expression of TF that converts the quiescent endothelium into a procoagulant surface. Increased thrombin generation at sites of vascular injury is believed to play a critical role in the pathogenesis of atherosclerosis. Antiplatelet therapy is commonly used in patients at risk of developing thrombotic disease. Dilazep is one of the antiplatelet drugs commonly used in European countries and Japan. Dilazep exerts antiplatelet activity by increasing the extracellular fluid level of adenosine in the systemic circulation by blocking adenosine uptake. Dilazep is also known to dilate coronary, cerebral, and renal vessels by blocking the calcium influx and to prevent the occurrence of thrombus formation in experimental animals. In preliminary clinical studies, we found that dilazep decreases the plasma level of various hemostatic markers (TAT, FDP-D-dimer, and fibrinogen) in patients with hypercoagulable state. Dilazep was also found to decrease the level of plasma TF antigen that is known to be a marker of endothelial cell or leukocyte activation. These findings prompted us to investigate the effect of dilazep on the procoagulant activity of HUVECs stimulated with TNF, thrombin, or PMA and on that of monocytes stimulated with TNF. The results of the present study showed that dilazep significantly inhibits the expression of TF in HUVECs and monocytes.

TF expression is known to be regulated at both transcriptional and posttranscriptional levels. Analysis of the TF mRNA expression showed that the thrombin- or PMA-induced TF expression in HUVECs is blocked by dilazep probably due to an effect on TF gene transcription and/or TF mRNA stability. PMA and thrombin are well-known activators of protein kinase C. This can be activated directly by PMA or indirectly by thrombin via the G protein-coupled receptor. Thus, dilazep probably...
The effect of dilazep on TF mRNA expression in HUVECs stimulated with TNF, thrombin, or PMA. (A) HUVECs were stimulated with each stimulant: 1,000 U/mL TNF, 25 nmol/L thrombin, or 5 nmol/L PMA for 3 hours in the presence (+) or absence (−) of 100 μg/mL dilazep. TF mRNA expression was analyzed using the RT-PCR method as described in the Materials and Methods. The amplified cDNA was blotted onto a nitrocellulose membrane and hybridized with the [32P]TF cDNA probe and then analyzed using an autoradiogram. (B) The data in (A) are depicted after normalization to equal amount of RNA loads based on the intensity of GAPDH mRNA. The data are expressed as the percentage of the control (each stimulant induced TF mRNA in the absence of dilazep). The values represent the mean ± SD of four independent experiments. Statistical analysis was performed using the Student’s t-test (*P < .05).

Inhibits the PMA- or thrombin-induced TF mRNA expression by blocking the protein kinase C-mediated signaling pathway. On the contrary, dilazep inhibited the HUVEC surface TF expression but it did not inhibit the expressions of total TF antigen and TF mRNA induced by TNF in HUVECs. This latter finding may be explained by the different signaling pathway of TNF-induced TF expression from that of thrombin- or PMA-induced TF expression. In this regard, TNF was previously found to induce TF mRNA expression in HUVECs by the sphingomyelinase/ceramide pathway.
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Fig 9. Effect of 8-SPT on the dilazep-induced downregulation of TF activity on the surface of HUVECs stimulated with TNF, thrombin, or PMA. HUVECs (2 x 10^5 cells/well) were incubated with (1 mmol/L, final concentration) or without (h) 8-SPT for 15 minutes and then with 100 μg/mL dilazep. After incubation of the cells with 1,000 U/mL TNF, 25 mmol/L thrombin, or 5 mmol/L PMA, in the presence or absence of dilazep, the cells were washed twice with 1 mL of HEPES-buffered saline containing 5 mmol/L CaCl_2. TF activity was determined as factor Xa generation on the surface of HUVECs as described in the Materials and Methods. The data are expressed as the percentage of the control (each stimulant induced TF activity in the absence of dilazep). Statistical analysis was performed using the Student's t-test (*P < .05; **P < .001).

Fig 10. The effect of varying concentrations of adenosine on TF activity expression on HUVECs stimulated with TNF, thrombin, or PMA. HUVECs were treated for 5 hours with each stimulant: 1,000 U/mL TNF (●), 25 mmol/L thrombin (■), or 5 mmol/L PMA (▲) in the presence of varying concentrations of adenosine (0, 2.5, 5, and 10 μmol/L). TF activity on HUVEC surface was determined as factor Xa generation using S2222 (see the Materials and Methods). The data are expressed as the percentage of the control (each stimulant induced TF activity in the absence of adenosine). The values represent the mean ± SD of four independent experiments. Statistical analysis was performed using the Student’s t-test (*P < .05; **P < .001).

In addition, TNF-mediated activation of protein kinase C is in general not required for nuclear factor-κB activation or for protein expression in endothelial cells.43,44 On the other hand, the fact that dilazep inhibited HUVEC surface expression of TF (decrease in TF activity and in its surface antigen expression by factor Xa generation assay and FACScan, respectively) without blocking the expression of total TF antigen and TF mRNA in these cells, as measured by ELISA and RT-PCR, suggests that dilazep regulates TNF-induced TF surface expression at the posttranscriptional level. Alternation in the folding, processing, and translocation of TF may be the potential mechanisms by which dilazep exerts this effect. Another piece of evidence that suggests the posttranscriptional regulation of TNF-induced TF expression by HUVECs is the results of the Western blot analysis showing that dilazep reduced the accumulation of matured forms (47 kD) of TF but increased the formation of TF with lower molecular weight (33 kD) in TNF-treated HUVECs. Similar lower molecular weight TF formation has been previously reported in lipopolysaccharide-treated mononuclear cells by endogenous cellular antioxidant potentiation.25 Dilazep is also known to act as an antioxidant agent and to modulate cytoplasmic redox state on endothelial cells.17 The redox state regulates protein synthesis at posttranscriptional level.
was determined as PCA as described in the Materials and Methods. The endothelial cell or monocyte surface is believed to play an essential role in the pathogenesis of atherosclerosis and thromboembolic complications. In the current study, we evaluated whether adenosine levels. In the current study, we evaluated whether adenosine potentiating activity of dilazep downregulates TF expression in HUVECs. The inhibitory effect of dilazep on TF expression in HUVECs stimulated with TNF, thrombin, or PMA was counteracted by adenosine receptor antagonists, suggesting that the inhibitory effect of dilazep is mediated, at least in part, by adenosine receptors. Adenosine itself was also found to inhibit TF expression in HUVECs. All these findings suggest that adenosine and its receptor may play important roles in the regulation of TF expression on endothelial cells.

In summary, this study showed that dilazep inhibits endothelial TF expression at both transcriptional and posttranscriptional levels and that its inhibitory effect is mediated, at least in part, by adenosine receptors. The results of this study also suggest that dilazep, besides its well-known antiplatelet effect, exerts a potent inhibitory activity on the procoagulant activity of endothelial cells and monocytes observed in these disorders.

The values represent the mean ± SD of four independent experiments.

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