Statins Enhance Migratory Capacity by Upregulation of the Telomere Repeat-Binding Factor TRF2 in Endothelial Progenitor Cells

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Background—Cultivation of endothelial progenitor cells (EPCs) leads to premature replicative senescence, limiting ex vivo expansion for potential clinical cell therapy. Recent studies have linked senescence to the dysfunction of telomeres, the “ends” of chromosomes, via the so-called mitotic clock or culture-induced stress. The purpose of this study was to elucidate a possible role of telomere biology in the functional augmentation of EPCs by statins.

Methods and Results—Human EPCs were isolated from peripheral blood. Using flow cytometry after fluorescence in situ hybridization with a telomere-specific (C3TA2)3 peptide nucleic acid probe (Flow-FISH), we found mean telomere length in untreated EPCs from healthy subjects to range between 8.5 ± 0.2 and 11.1 ± 0.5 kb with no change over 6 days of culture, excluding telomere erosion as one cause for premature senescence. Although mean telomere length did not differ between statin-treated and untreated EPCs, atorvastatin (0.1 μmol/L) and mevastatin (1.0 μmol/L) both led to a more than 3-fold increase in the expression of the telomere capping protein TRF2 (telomere repeat-binding factor), as shown by immunoblotting, whereas quantitative reverse transcription–polymerase chain reaction demonstrated no increase in TRF2 mRNA. Telomere dysfunction of EPCs was also paralleled by a 4-fold increase in the DNA damage checkpoint-kinase 2 (Chk2). Conversely, statin cotreatment or overexpression of TRF2 completely suppressed Chk2 induction. Finally, overexpression of a dominant negative mutant of the TRF2 protein abrogated statin-induced enhancement of migratory activity down to baseline values.

Conclusions—Ex vivo culturing of EPCs leads to “uncapping” of telomeres, indicated by the loss of TRF2. Statin cotreatment of EPCs prevents impairment of their functional capacity by a TRF2-dependent, posttranscriptional mechanism. This is the first time a beneficial effect of statins on telomere biology has been described. (Circulation. 2004;110:3136-3142.)

Key Words: telomere ■ aging ■ stem cells ■ telomere repeat-binding factor 2

Endothelial progenitor cells (EPCs) can be isolated from bone marrow, cord blood, and blood-derived mononuclear cells.1,2 The diversity of their potential therapeutic effects in the cardiovascular system comprises the enhancement of endothelial and ventricular function after myocardial infarction, the facilitation of angiogenesis by secretion of chemokines, and their antiatherogenic action.3–7 Unfortunately, the limited number and impaired migratory function of EPCs in patients with coronary artery disease offsets their full therapeutic potential.8 First, the relative paucity of the cells among white blood cells mandates ex vivo expansion as a prerequisite to clinical administration; second, we found migratory capacity to be an independent predictor of subsequent infarct size reduction after intracoronary infusion in patients with acute myocardial infarction.9 In previous studies, we found that HMG-CoA reductase inhibitors (statins) (1) delay replicative senescence in endothelial cells, (2) reduce premature senescence in cultured EPCs,10,11 and (3) improve migratory function,12 together leading to our routine use of statins in preparing EPCs for human infarct repair (TOP-CARE).4 Premature senescence of cultured cells can be triggered by oxidative stress, leading to telomere dysfunction.13,14 Telomeres consist of tandem T2AG3 repeats at the chromosome ends, maintained by the enzyme telomerase reverse transcriptase (TERT) and bound by specific telomere repeat-binding factors (TRFs), including TRF1 and TRF2.15–20 Of these, TRF1 functions at least in part as a negative regulator of telomere length. Telomere dysfunction can occur because of the loss of TRF2, implying a role of telomere “uncapping” rather than telomere shortening.21–27

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Here, we demonstrate that statins delay premature senescence in cultivated EPCs independently of mean telomere length. Statin-enhanced migratory activity of EPCs also depends in part on the induction of the telomere-capping protein TRF2. This is the first time a protective effect of statins on telomere biology has been described.

Methods

Materials

Human recombinant vascular endothelial growth factor (VEGF), macrophage-colony-stimulating factor, granulocyte-macrophage colony-stimulating factor, and interleukin-4 were purchased from Cell Concepts. Antennapedia internalization sequence peptide was synthesized by BioSynth.

Cell Culture

Mononuclear cells were isolated by density-gradient centrifugation with Ficoll from peripheral blood of healthy human volunteers as described previously.\(^{24}\) Immediately after isolation, total mononuclear cells (8×10\(^4\) cells/mL; cell density, 2.5×10\(^6\) cells/cm\(^2\)) were plated on culture dishes coated with human fibronectin (Sigma) and maintained in endothelial basal medium (EBM; Clonetics) supplemented with EGM SingleQuots, VEGF (100 ng/mL), and 20% FCS. More than 90% of adherent EPCs show endothelial characteristics, as demonstrated by Dil-acetylated LDL uptake and lectin binding and immunostaining of typical endothelial marker proteins, including VEGF receptor (VEGFR2) (KDR) (Reflia Tech), endoglin (CD105) (NeoMarkers), and von Willebrand factor (Oncogene). Using FACS analysis, we found 94% of EPCs to stain positive for KDR, 76% for CD105, and 72% for von Willebrand factor.

Q-FISH

Endothelial cells or EPCs were incubated for 24 hours with 100 ng/mL colcemide. Nuclei were isolated by hypotonic swelling in a 60-mmol/L KCl solution and fixed in a 3:1 solution of methanol/glacial acid. Hybridization was performed with a Cy3-conjugated (C\(_3\)TA\(_2\))\(_3\) peptide nucleic acid (PNA) probe (Dako) according to the manufacturer’s instructions. Counterstaining was performed with 4',6-diamino-2-phenylindole (DAPI; 2 \(\mu\)g/mL).

Determination of Human Telomere Length by Flow-FISH

The average length of telomere repeats at chromosome ends in individual peripheral blood leukocytes was measured by Flow-FISH as previously reported.\(^{29–32}\) EPCs were detached by trypsin, and the cell suspension was centrifuged for 5 minutes at 800 \(g\). The supernatant was removed, and the cell pellets were washed twice with PBS. A total of 2×10\(^5\) EPCs and 1×10\(^5\) calf thymocytes (as an internal calibration standard) were resuspended together in the hybridization mixture. After an initial reaction time of 10 minutes at room temperature, the cell pellets were washed with 1 mL washing buffer A (75% formamide/glacial acid). Hybridization was performed with a Cy3-conjugated (C\(_3\)TA\(_2\))\(_3\) peptide nucleic acid (PNA) probe (Dako) according to the manufacturer’s instructions. Counterstaining was performed with 4',6-diamino-2-phenylindole (DAPI; 2 \(\mu\)g/mL).

Subpopulation-specific telomere lengths were calculated by subtracting the mean fluorescence of the background control (no probe) from the mean fluorescence obtained from cells hybridized with the telomere probe. FITC-labeled fluorescent beads (QuantumTM-24 Premixed; Flow Cytometry Standards) were used to correct for daily shifts in the linearity of the flow cytometer and fluctuations in the laser intensity and alignment. Mean telomere length of EPCs was calculated as molecular equivalents of soluble fluorochrome (EPCs/molecular equivalents of soluble fluorochrome (thymocytes) times the mean telomere length of thymocytes determined directly by Southern blotting.

Whole-Cell Extracts

Cells were washed 3 times in cold PBS and then lysed for 30 minutes at 4°C in lysis buffer containing 20 mmol/L Tris-HCl (pH 7.4), 1 mmol/L EDTA (pH 8.0), 1 mmol/L EGTA, 150 mmol/L NaCl, 1% Triton, 2.5 mmol/L Na-pyrophosphate, 1 mmol/L \(\beta\)-glycerophosphate, 1 mmol/L Na-orthovanadate, and the following protease inhibitors: 200 mmol/L PMSE and 1 \(\mu\)g/mL leupeptin. After centrifugation at high speed, the supernatant was collected and the protein content of all samples was determined using the Bio-Rad protein assay with \(\gamma\)-globulin as a standard.

Western Blot Analysis

Electrophoresis was performed on 12% SDS-polyacrylamide gels loading 30 \(\mu\)g of protein per lane. After transfer to a 0.2-\(\mu\)m polyvinylidene difluoride membrane (Bio-Rad), the membranes were blocked in 3% BSA (in PBS, pH 7.5, and 0.2% Tween 20) for 1 hour and incubated overnight with the indicated primary antibody (Ab) at 4°C. Each Ab incubation period was followed by 1 hour of membrane washing in 0.1% Tween-20 in PBS. Detection was performed by use of a horseradish peroxidase–linked anti-rabbit (1:7500 in PBS/1% Tween 20/2% nonfat milk), anti-goat (1:5000), or anti-mouse (1:5000 in PBS/1% Tween 20/2% nonfat milk) (secondary) Ab (Amersham) and the Amersham enhanced chemiluminescence system (ECL kit). The primary Abs were used: \(\beta\)-actin mouse monoclonal Ab (Sigma A5441; 1:8000), TRF2 H-300 rabbit polyclonal Ab (Santa Cruz sc-9143; 1:250), TRF2 goat polyclonal Ab (Santa Cruz sc-8528; 1:250), Checkpoint-kinase 2 (Chk2) mouse monoclonal Ab (Santa Cruz sc-17747; 1:250), green fluorescent protein (GFP) rabbit polyclonal Ab (Invitrogen 46-0092; 1:3000), and c-nect 9B11 mouse monoclonal Ab (Cell Signaling 2276; 1:500).

Quantitative Real-Time Reverse Transcription–Polymerase Chain Reaction

Total RNA from 10\(^5\) EPCs was isolated from the guanidine method using Qiagen’s RNA extraction kit. Quantification of TRF2 mRNA was performed by amplification of EPC cDNA in a 1-step reverse transcription–polymerase chain reaction (RT-PCR) using the LightCycler (Roche Diagnostics) real-time thermocycler according to the instructions of the manufacturer (Roche Diagnostics). The RT-PCR reaction was composed of 3.25 mmol/L Mn(OAc)\(_2\), 1× LightCycler–RNA Master SYBR Green I, 100 ng mRNA, and 0.5 mmol/L of each primer in a final volume of 20 \(\mu\)L. The following oligonucleotide primers were used for amplification of human TRF2 cDNA: sense (5’→3’) CCCAAGAACAAGCGCATGAC and antisense (5’→3’) TTTCTGCACTCCAGCCTTGAC. Amplification was performed with 45 cycles and an annealing temperature of 61°C. Copy numbers were calculated by the instrument software (Roche Diagnostics) from standard curves of an in vitro–transcribed cytokine mRNA. The specificity of the amplification reaction was determined by a melting curve analysis. PCR products were analyzed on a 2% agarose gel to confirm a 341-bp product.

Adenoviral Constructs and Infection of EPCs

Plasmids for human TRF2 and its dominant-negative truncation (TRF2\(_{1893}\)) were provided by T. de Lange (Rockefeller University, New York, NY, USA). Adenoviruses coexpressing enhanced GFP were generated by use of pAdTrack-cytomegalovirus and...
pShuttle-cytomegalovirus (provided by B. Vogelstein, Johns Hopkins Oncology Center, Baltimore, Md 34,35) and have been described previously. 24 If not otherwise stated, Antennapedia peptide (Antennapedia internalization sequence RQIAIWFQNR-RMKWAA36) at a concentration of 0.5 mmol/L, purified adenovirus, and 100\%/H9262L of OptiMem (Gibco) were incubated for 30 minutes at room temperature. The mixture was then added to freshly isolated mononuclear cells (see Cell Culture) in a 2.5-mL volume and left together with the cells until day 3. Adenoviral infection was validated by visualization of enhanced GFP under fluorescence microscopy.

**Assessment of Migratory Capacity of EPCs**

After 3 days of culture, progenitor cells were resuspended in EBM without FCS, containing 1% BSA. Then 5×10^5 EPCs were placed in the upper chamber of a modified Boyden chamber (Falcon HTS Fluoro Blok insert, No. 351152, 8-μm pore size). The chamber was placed in a 24-well culture dish (Falcon, No. 353504) containing endothelial basal medium, 50 ng/mL VEGF, and 100 ng/mL stromal cell–derived factor (SDF)-1 (where indicated) for measuring the migratory capacity of EPCs. After 24 hours of incubation at 37°C, the lower side of the filter was washed with PBS and fixed with 2% paraformaldehyde. For quantification, cells were stained with Dil-acetylated LDL and were counted manually in 3 random microscopic fields per well.

**Results**

**Lack of Telomere Erosion in Senescent EPCs**

To address the possibility of telomere erosion as a cause of premature senescence and dysfunction in cultured EPCs, we performed the flow cytometric determination of the mean telomere length after fluorescence in situ hybridization (Flow-FISH) with labeled telomere probes as described previously.29–32,37 In situ telomere hybridization (Q-FISH) was performed to prove the specificity of the used (C3TA2)3 PNA probe in EPCs (data not shown). EPCs were analyzed in interphase. Using Flow-FISH, we found the mean telomere length in EPCs cultured from 3 different donors. Mononuclear cells were incubated for 3 days either with DMSO (0.1%; Cont) or atorvastatin (AT; 0.1 μmol/L). Data are mean±SD, n=3. D, Telomere length of EPCs on day 4 and day 6 after incubation of mononuclear cells with either DMSO (0.1%; Cont) or atorvastatin (AT; 0.1 μmol/L). Data are mean±SEM, n=3.
length by itself can conceal small populations of critically shortened telomeres, we also specifically compared the potential effects of propagation and statins on the shortest length of telomeres in the Flow-FISH distribution and excluded a shift in the length distribution. Together, these findings would better support the concept of telomere dysfunction independent of critical telomere shortening, such as because of telomere uncapping.

Statins Induce TRF2 by a Posttranscriptional Mechanism

To investigate the possibility that statins might, instead, have salutary effects on telomere-associated proteins in cultured EPCs, we first examined the protein expression of the specific TRFs. Although expression of TRF1 did not change under statin treatment (data not shown), TRF2 was induced more than 3-fold by atorvastatin or mevastatin, in a dose-dependent manner, after 3 days of cultivation (Figure 2, A and B), and induction reached its maximum at 0.1 μmol/L and 1.0 μmol/L, respectively. Simvastatin (data not shown) had an identical effect on TRF2 expression at a concentration of ≥0.5 μmol/L. To obtain this effect, which was consistent among each of 3 healthy blood donors (Figure 2, A and B), statin incubation had to be initiated in the first 24 hours after isolation of mononuclear cells (Figure 2C). Addition of atorvastatin at later time points did not lead to an increase of TRF2 expression.

We previously showed that statins inhibit premature senescence in EPCs by preventing formation of mevalonate and the downstream products farnesylpyrophosphate (FPP) and geranylgeranylpyrophosphate (GPP).11 As shown in Figure 2D, mevalonate, FPP, and GPP partially repressed TRF2 induction by atorvastatin, indicating a mechanism involving isoprenylation. To test for induction of TRF2, we performed semiquantitative PCR using TRF2-specific oligonucleotide primers (Figure 2E). However, no statistical difference in TRF2 mRNA expression could be seen among the statin-treated and untreated cells, suggesting an enhancement of protein translation or stability rather than mRNA induction.

Telomere Dysfunction of EPCs Leads to Chk2 Induction

Because loss of the telomere-capping protein TRF2 leads to telomere dysfunction, as measured by activation of the DNA damage protein kinase pathway, we investigated this possible

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**Figure 2.** Induction of TRF2 protein expression by statins. A, Mononuclear cells were isolated from healthy donors, and whole protein extracts were generated on day 3 from EPCs. Representative immunoblots from 3 different donors are shown, using Abs against TRF2 (H-300), Chk2, and β-actin. Cells were incubated either with 0.1% DMSO (Co), atorvastatin (At; 0.1 μmol/L), or mevastatin (Me; 1.0 μmol/L). B, Semiquantitative analysis of TRF2 immunoblot (A) by densitometry. Control value is defined as 100%. All values are presented as mean±SEM. §P<0.01, #P<0.001 vs control. C, Mononuclear cells were isolated from healthy donors, and whole protein extracts were generated on day 6 from EPCs, EPCs were treated either without (-) or with 0.1 μmol/L atorvastatin, added at indicated time points, where, only in lane marked with an asterisk (*), atorvastatin was already discontinued after 72 hours. D, Mononuclear cells were cultivated for 4 days in EBM with supplements and 20% FCS in presence or absence of atorvastatin (0.1 μmol/L), mevalonate (Mev; 200 μmol/L), farnesylpyrophosphate (FPP; 10 μmol/L), and geranylgeranylpyrophosphate (GPP; 10 μmol/L). E, Semiquantitative RT-PCR of TRF2 mRNA in EPCs using Lightcycler (Roche). Freshly isolated mononuclear cells were cultivated in EBM with supplements and 20% FCS in presence or absence of atorvastatin (AT; 0.1 μmol/L). DMSO (0.1%) served as a vehicle control. mRNA was isolated at 0 hours from mononuclear cells or after 24 and 48 hours from adherent cells. Data are mean±SEM, n=2. One of 3 representative experiments is shown.
downstream contender in TRF2 signaling. Activity of Chk2, a DNA damage kinase, increases accompanying partial loss of TRF2 in human myocardium during heart failure and is triggered in rat cardiomyocytes by interference with TRF2 function or expression. Indeed, Chk2 protein level in EPCs increases under culture conditions to more than 4-fold compared with levels in freshly isolated monocytes (Figure 3A), whereas atorvastatin, mevastatin, or simvastatin (data not shown) cotreatment completely abrogated Chk2 induction (23% versus 100% control; Figures 2A and 3B). Finally, overexpression of TRF2 wild-type protein in EPCs also prevented Chk2 upregulation (Figure 3C), suggesting that Chk2 is a downstream target of telomere dysfunction induced by lack of TRF2 in EPCs.

Enhancement of Migratory Activity by Statins Depends on TRF2 Expression

Finally, to provide evidence that the increase in TRF2 contributes to the increased functional capacity of EPCs elicited by statins, we expressed dominant-negative TRF2 in EPCs. To achieve this, freshly isolated mononuclear cells were infected with adenoviral constructs. The validity of viral vectors was confirmed by immunoblots in the day 3 EPCs (Figure 4A). Transduced EPCs were used on day 3 for migration assays, showing an increase in basal migratory activity after coculture with 0.1 μmol/L atorvastatin (AT; 0.1 μmol/L) was added to media where indicated. Migratory activity of EPCs was determined on day 3 after adenoviral infection either without (B) or with (C) addition of 100 ng/mL SDF (P<0.05; §P<0.01; #P<0.001).

Figure 3. Regulation of Chk2 in cultured EPCs. A, Immunoblot showing upregulation of Chk2 in EPCs, cultured for 3 days without addition of atorvastatin (AT; 0.1 μmol/L). Freshly isolated mononuclear cells were used as a control (Co) for day 0. A representative blot from 2 donors is shown. B, Semiquantitative analysis of Chk2 immunoblot (Figure 2A) by densitometry. Control value is defined as 100%. All values are presented as mean±SEM. #P<0.001 vs control. C, Overexpression of wild-type TRF2 in freshly isolated mononuclear cells abrogated induction of Chk2 in day 3 EPCs. Increasing concentrations of Antennapedia peptide (Antp) were used during adenoviral infection as indicated. A representative blot from 2 experiments is shown.

Figure 4. Statin-enhanced migration in EPCs depends on TRF2 function. A, Viral vector. Dominant-negative TRF2 (TRF2 B M) tagged with myc epitope lacks Myb domain and N-terminal basic domain, coexpressing enhanced GFP. Western blots confirming expression of exogenous proteins in EPCs. Dominant negative TRF2 is detected with H-300 Ab (raised against amino acids 49 to 300; Santa Cruz Biotechnology) but not C-16 (raised against C terminus, data not shown). B and C, Freshly isolated mononuclear cells were infected with adenoviral vectors Ad-GFP (GFP) and Ad-TRF2dn (dn), respectively. Atorvastatin (AT; 0.1 μmol/L) was added to media where indicated. Migratory activity of EPCs was determined on day 3 after adenoviral infection either without (B) or with (C) addition of 100 ng/mL SDF into lower chamber. Number of Dil-positive cells/HPF was counted. Data are mean±SEM. Three parallel experiments with n=3 HPFs each were included in analysis. Levels of significance between pairs of groups in B and C are indicated by following symbols: *P<0.05; §P<0.01; #P<0.001.
EPCs expressing dominant-negative TRF2 was reduced almost down to baseline (no statin pretreatment): 18.8±1.9 cells/HPF without SDF (P<0.001 versus Ad-GFP+atorvastatin; Figure 4B) and 53.7±3.9 cells/HPF with SDF attraction (P<0.01 versus Ad-GFP+atorvastatin; Figure 4C), respectively.

**Discussion**

The results of the present study demonstrate that statins induce expression of a telomere repeat-binding factor, TRF2, in cultured EPCs, thereby improving functional capacity and providing a potential explanation for the effect of statins on cell senescence.

Human telomeres are bound by 2 double-strand telomere binding proteins, TRF1 and TRF2, each of which recruits another set of interacting proteins. Removal of TRF2 triggers apoptosis or senescence and can result in fusion of the chromosome ends. Loss of TRF2, however, occurs in end-stage human heart failure and leads to telomere dysfunction. Overexpression of TRF2 and TERT is protective against oxidative stress in cultured cardiomyocytes. Similarly, we demonstrate that interference with TRF2 abolishes the statin-induced enhancement of the migratory capacity in EPCs. Of note, the migratory activity of EPCs reflects the most relevant parameter with regard to functional improvement after cell therapy for acute myocardial infarction. It is also possible that the induction of TRF2 by statins leads to “juvenation” of EPCs, because replicative senescence is delayed. Younger cells would migrate faster than older cells because of their age characteristics. In this case, TRF2 protein might correlate very well with cellular aging, thus exerting only an indirect role on cell migration.

Telomere length in human white blood cells shortens during aging. Interestingly, individuals with shorter white blood cell telomeres after adjustment for age show a 3-fold higher mortality rate from heart disease and more than 8-fold increase in mean telomere length occurred as an explanation of how statins favorably affect senescence in human EPCs. Of note, the migratory activity of EPCs reflects the most relevant parameter with regard to functional improvement after cell therapy for acute myocardial infarction. It is also possible that the induction of TRF2 by statins leads to “juvenation” of EPCs, because replicative senescence is delayed. Younger cells would migrate faster than older cells because of their age characteristics. In this case, TRF2 protein might correlate very well with cellular aging, thus exerting only an indirect role on cell migration.

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