High-level expression of Egr-1 and Egr-1–inducible genes in mouse and human atherosclerosis

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To understand the mRNA transcript profile in the human atherosclerotic lesion, RNA was prepared from the fibrous cap versus adjacent media of 13 patients undergoing carotid endarterectomy. cDNA expression arrays bearing 588 known genes indicated that lesions express unexpectedly high levels of the early growth response gene, Egr-1 (NGFI-A), a zinc-finger transcription factor that modulates a cluster of stress-responsive genes including PDGF and TGF-β. Expression of Egr-1 was an average of 5-fold higher in the lesion than in the adjacent media, a result confirmed by RT-PCR, and many Egr-1–inducible genes were also strongly elevated in the lesion. Time-course analyses revealed that Egr-1 was not induced ex vivo. Immunocytochemistry indicated that Egr-1 was expressed prominently in the smooth muscle–actin positive cells, particularly in areas of macrophage infiltration, and in other cell types, including endothelial cells. Induction of atherosclerosis in LDL receptor-null mice by feeding them a high-fat diet resulted in a progressive increase in Egr-1 expression in the aorta. Thus, induction of Egr-1 by atherogenic factors may be a key step in coordinating the cellular events that result in vascular lesions.

1–mediated gene transcription plays a key role in orchestrating the functional characteristics of the vessel wall after arterial injury.

Atherosclerosis is believed currently to be an excessive fibroproliferative response to chronic vascular injury by diverse agents. The pioneering work of Dr. Russell Ross, and many others, described the involvement of peptide growth factors and cytokines, such as PDGF and FGF, in orchestrating migration, proliferation, and matrix production by SMC-like cells, macrophages, lymphocytes, and endothelial cells (19). Other platelet-derived factors, such as TGF-β, act as autocrine/paracrine cellular products with multiple effects on proliferation, migration, and matrix synthesis. The coordinate regulation of these growth factor and cytokine signals is probably a key determinant of the cellular events that occur during atherogenesis. This regulation may be mediated by alterations in the amount or localization of various transcription factors within the lesion cells.

In the course of screening the mRNA expression pattern of human atherosclerotic lesions using cDNA arrays, it was observed that Egr-1 levels often exceeded the levels of constitutive genes, such as actin, and that Egr-1 transcript levels were much higher in the lesion than the adjacent media. The present studies examined the expression of Egr-1– and Egr-1–responsive genes in these lesions. Elevated levels of Egr-1 were associated with increases in Egr-1 target genes. Immunocytochemistry was used to determine the expression pattern in the human carotid atherosclerotic lesion. Atherosclerosis in the LDL receptor–null mouse was also associated with increased Egr-1 expression. Collectively, these results implicate Egr-1 induction, and subsequent target gene expression, in human and mouse atherogenesis.

**Methods**

**Tissue specimens.** Vascular specimens were obtained during surgical revascularization procedures at The New York Presbyterian/Cornell Medical Center (New York, New York, USA) as waste surgical specimens under institutional review board–approved protocols. Surgical endarterectomy of carotid artery disease produces full diameter lesions of 2–5 cm in length that commonly contain tunica media, without adventitia. The medial tissue can be cleanly dissected from the overlying lesion. Internal mammary arteries were obtained as excess waste from coronary artery bypass grafting (CABG) operations.

**Timeline of postsurgical sample processing.** Under normal conditions, the specimens could be obtained in saline, returned to the laboratory, dissected under sterile conditions, and placed into RNAzol within 40 to 60 minutes of surgical excision. To investigate the possibility that early response genes were induced ex vivo within this time frame, dissection of the lesion was conducted within the surgical suite, and the specimens could be placed into RNAzol within 5 minutes. Parallel portions of tissue were then held in saline for 1 hour before being placed in RNAzol.

**Immunocytochemistry.** Portions of endarterectomy specimens were equilibrated with 30% sucrose and then embedded and sectioned in OCT. Frozen sections were immunostained with an Egr-1–specific antibody (C-19; Santa Cruz Biotechnology, Inc., Santa Cruz, California, USA) at a 1:50 dilution by standard procedures for the avidin-biotin–enhanced immunoperoxidase detection (Vector Laboratories, Burlingame, California, USA). Positive immunostaining was detected with diaminobenzidine tetrahydrochloride (DAB) and counterstained with hematoxylin.

In some patients (n = 4), sections were double-stained with rabbit polyclonal antibodies to Egr-1, detected by avidin-biotin peroxidase and DAB substrate (brown), and counterstained with mAbs to marker antigens detected with alkaline phosphatase–labeled secondary antibodies and NBT/BCIP substrate, producing a blue product. Cross-reactivity of the secondary antibodies (anti-rabbit and anti-mouse) was excluded with controls that employed only 1 antibody and a nonimmune antibody of the other species. Cell type-specific antibodies included SMC–specific α-actin (Clone 1A4; Sigma Chemical Company, St. Louis, Missouri, USA) and macrophage-specific (HAM56; DAKO Corp. Carpinteria, California, USA) and lymphocyte-specific leukocyte common antigen (LCA) (CD45; DAKO).

**RNA purification.** Tissues were transiently immersed (3 minutes) in RNAlater (Ambion Inc., Austin, Texas, USA) to minimize degradation and stabilize the RNA. Total RNA was purified from vascular specimens or LDC using RNAzol with minor modifications of the manufacturer’s method. In particular, early precipitation reactions could not be performed on ice, as specified, because of excessive precipitation of polysaccharides from the lesions. RNA quantity and quality was assessed by agarose gel electrophoresis and ethidium bromide staining.

**RT-PCR.** RNA (1–5 μg) was reversed transcribed with Moloney murine leukemia virus (MMLV) reverse transcriptase (Perkin-Elmer, Foster City, USA) using random hexamer primers. A portion of this cDNA (250–500 ng) was then used for PCR with Taq (GeneAmp; Perkin-Elmer) under the following conditions (95°C denaturing temperature for 1 minute, 60°C anneal/extend for 25–40 cycles, with a final extension of 72°C for 10 minutes). Primers for Egr-1 were directed at 1847–1868 and 2371–2392 of the cDNA (GenBank M62829), producing a 545-bp product. Human β-actin mRNA was detected with primers directed at 484–504 and 813–833 (GenBank AB004047), producing a 350-bp product. Human RhoA (GenBank AB004047), producing a 350-bp product. Human RhoA (GenBank L25080) was detected with primers annealing to 290–317 and 545–572, producing a 283-bp product.

**Genomic expression arrays.** For genomic analysis, RNA was further purified by retention on glass fiber columns (HighPure; Boehringer-Mannheim, Indianapolis, Indiana, USA), DNAse digestion, ethanol precipitation, and a final gel filtration on a desalting column (Chroma-spin 200 CLONTECH Laboratories Inc., Palo Alto, California, USA) to remove small
oligonucleotides or contaminants. The paired RNAs (i.e., media versus lesion) were reversed transcribed with MMLV, $^{32}$P-ATP, and a mixture of 588 sequence-specific primers, as described by the manufacturer (CLONTECH). In some studies, the reverse transcription–labeling step was performed with Superscript (GIBCO BRL, Gaithersburg, Maryland, USA), an RNase H-MMLV mutant that appeared to produce more efficient and consistent labeling. The labeled cDNAs were desalted to remove free isotope and then hybridized overnight to identical cDNA arrays under stringent conditions (68°C). The array membranes were washed with 2 $\times$ SSC with 1% SDS for 30 minutes, 4 times, and 0.1 $\times$ SSC with 0.5% SDS for 30 minutes, 3 times, all at 68°C. The paired membranes were exposed to storage phosphor plates (Eastman Kodak Co. Scientific Imaging Systems, New Haven, Connecticut, USA) for 1–5 days and quantified on a phosphoimager (Storm; Molecular Dynamics, Sunnyvale, California, USA).

Mice, diet, and immunohistochemical analysis. Low density lipoprotein (LDL) receptor-negative (LDLR–/–) mice were created by homologous recombination as described (20). Mice were placed on a high-cholesterol diet consisting of 1.25% cholesterol (including 7.5% cocoa butter, 7.5% casein, and 0.5% cholic acid). Mice were sacrificed after 0, 5, 10, and 20 weeks on the diet. These time points were selected based on preliminary studies that demonstrated lesions begin to develop in the aortic arch and near the ostia after 4 weeks on the diet. After sacrifice, the arterial tree was perfused with PBS (40 mL/10 min) and then perfusion-fixed with buffered formaldehyde (70 mL/15–20 min) at a pressure of 100 mmHg. Samples of the aorta were embedded in OCT and immunostained with the Egr-1 antibody as described above, using a fast red substrate for the peroxidase-labeled secondary antibody. Sections were lightly counterstained with methylene green.

Results

Expression of Egr-1 mRNA and related genes in human atherosclerotic carotid arteries: Analysis by genomic expression arrays. To understand the physiological and pathological state of the cells that compose the fibroproliferative component of the atherosclerotic lesion, tissues from the fibrous cap of lesions were compared with the adjacent, underlying tunica media of 13 patients with obstructive carotid atherosclerosis. RNA was simultaneously reverse transcribed, labeled radioactively, and then hybridized to spatially addressable cDNA arrays (Figure 1a). The results were quantified by Storm phosphoimager, the background subtracted, and the counts normalized by an average of 3 housekeeping genes (GAPDH, $\beta$-actin, and 23-kDa highly basic protein; Figure 1b).

The overall results of the studies showed that the level of gene expression in lesion tissue was significantly correlated with expression in the tunica media ($r = 0.87, P < 0.05$), an outcome consistent with both tissues possessing many similar basic functions. Quantitation of the arrays proved accurate and reliable: re-exposure of the same cDNA membrane for 1 day versus 4 days, followed by quantitation of each membrane, revealed a strong positive correlation ($r = 0.9991, P < 0.001$).

One surprising outcome from this genomic analysis was the relatively high-level expression of the Egr-1 mRNA transcript, a gene previously thought to be associated with the early growth response. Egr-1 mRNA levels were extremely high in the atherosclerotic lesion: on average they were 2-fold higher (200%) than the housekeeping genes ($n = 13$ patients, 3 experiments, range = 101–285%). Levels were consistently lower in the adjacent media: an average of only 42% of housekeeping genes (range 9–71%). Thus, there was an almost 5-fold induction of Egr-1 expression in the atherosclerotic lesion (Figure 1b).

![Figure 1](https://example.com/figure1.png)

**Figure 1**

Analysis of mRNA levels in human atherosclerotic lesions by cDNA arrays. The mRNA was purified from the fibrous cap (lesion) or the tunica media (media) of 13 patients and analyzed in pooled groups by hybridization to cDNA arrays. (a) Storage phosphor image of 1 array containing 588 unique cDNAs double-spotted onto nitrocellulose and hybridized with $^{32}$P-labeled, reverse-transcribed cDNA from human media acquired by endarterectomy of 6 pooled patients. (b) Quantitation from the storage phosphor screen was normalized to a mean housekeeping value (mean), which was the average of GAPDH, $\beta$-actin, and 23-kD HBP for each array. Thirteen patients were analyzed in groups on 3 array pairs (media versus lesion). Error bars, SEM; $n = 3$ experiments. HBP, highly basic protein.
Figure 2
Analysis of mRNA levels in human lesion by RT-PCR. (a) The mRNA was prepared from the dissected fibrous cap of the lesion (L) or the adjacent tunica media (M) within 5 minutes of surgical excision or after allowing an identical portion to incubate under transfer conditions for 65 minutes. The mRNAs were then analyzed by RT-PCR with primers directed at Egr-1 or RhoA. Two patients (E213, E217) are shown, demonstrating the difference noted in 3 of the 4 patients analyzed. (b) The difference in mRNA levels between media and lesion tissue was semiquantitated by serial dilution of the L mRNA before RT-PCR for Egr-1 (upper band) or β-actin (lower band). Nonspecific amplification or contamination were not detected in water (H2O) control.

By comparison, other, similar transcription factors were expressed at lower levels and were not increased in lesions. The transcription factor CREB showed only a small increase from 3 to 4%. The mRNA levels of the Sp1 transcription factor, an important competitor for Egr-1 at certain promoter domains (4, 5), was elevated from 2% in medial tissues to 13% in lesion tissue. Egr-1 is the prototype member of the NGFI-A (nerve growth factor inducible-type A) family of transcription factors. The NGFI-B family of factors may also have transcription factor activity, but includes nuclear orphan receptors similar to thyroid and steroid receptors. This NGFI-B family includes the TR3 factor, which was contained within the same arrays and was expressed at low levels that did not differ between lesion and media (Figure 1b).

Confirmation of Egr-1 mRNA expression by RT-PCR analysis. Because the cDNA arrays depend upon hybridization of labeled cDNA reverse transcripts to immobilized capture probes (averaging 300 nucleotides), the potential for nonspecific hybridization exists. Whereas the high stringency of the hybridization and washing should suppress nonspecific hybridization, the mRNA from 4 additional patients, not included in the array analysis, was prepared under highly controlled conditions and analyzed by RT-PCR. Consistently higher levels of amplimers for Egr-1 (550 bp) were observed in the lesion (L) than in the adjacent media (M) of the same patient (Figure 2a, 2 patients shown). The amount of mRNA used for amplification was identical in both tissues, a fact confirmed by a separate RT-PCR for RhoA. The difference in Egr-1 mRNA levels was semiquantitated in these samples by serial dilution of the lesion (L) sample until it reached a level similar to the media (M). As shown in Figure 2b, the E213 lesion sample could be diluted 4-fold before it was comparable to the media levels, and the E217 sample could be diluted 8-fold before comparable levels were achieved, thus supporting the 5-fold difference observed in the arrays. The Egr-1 amplimer was sequenced with 4-color dye-terminator methods and found to match identically with the Egr-1 cDNA sequence (M62829) over 300 bp of the accurately readable region. These studies demonstrate that Egr-1 transcript expression was detected at a 5-fold higher level in lesions than in the adjacent media and confirm the findings from the genomic expression arrays.

Effect of ex vivo duration on Egr-1 expression. Because Egr-1 is an early response gene, the induction of Egr-1 expression could have occurred ex vivo, as a result of stress resulting from vascular surgery and postoperative dissection. To determine whether the interval between surgical excision of the specimen and the preparation of RNA affected Egr-1 induction, patient specimens were prepared under conditions that allowed the rapid dissection and fixation of the tissue (5 minutes) whereas identical portions of tissue were allowed to incubate under conditions similar to the array experiments (maximally 65 minutes). The results are shown in Figure 2a, which indicates that regardless of whether the RNA was derived from lesion (L) or media (M), Egr-1 levels did not increase in the ex vivo conditions used to transport the specimens from the surgical suite to the laboratory. Rather, Egr-1 levels tended to decrease slightly, though the differences between media and lesion remain. Furthermore, if the Egr-1 levels were induced ex vivo, then both the lesion and media tissues should be induced similarly, which is not observed.

Expression of Egr-1 mRNA in lesions versus internal mammary arteries. The medial portions of the endarterectomy specimens are a useful control for the adjacent atherosclerotic lesion, but typically are not histologically

Figure 3
Analysis of mRNA for Egr-1 in IMA versus atherosclerotic carotid arteries. IMA were acquired from patients undergoing CABG. The mRNA was prepared from tunica media of 4 patients (31–34) and, for reference, compared with reamplification of E217 media and lesion, as shown in Figure 2. Primers were directed at Egr-1 or β-actin for normalization. Nonspecific amplification or contamination were not detected in water (H2O) control.
normal vessels by virtue of their proximity to an occlusive, fibroproliferative lesion. To determine Egr-1 expression in a less-diseased vessel, normal internal mammary arteries (IMA) were acquired from CABG, and RNA was prepared as rapidly as possible given different surgical conditions (average 15 minutes). RNAs from 4 patients were analyzed by RT-PCR for Egr-1 and compared with RNA from E217 media and lesion (Figure 2), which was reamplified under identical conditions. As shown in Figure 3, E217 lesion RNA again showed higher levels of Egr-1 expression than the adjacent media, and levels in the IMA ranged from undetectable to just higher than the E217 media. The presence of amplifiable mRNA was confirmed by parallel PCR reactions for β-actin. Thus, there appears to be less Egr-1 transcript in normal IMA than in vascular lesions.

Expression of Egr-1 antigen in human tissues by immunocytochemistry. To determine the pattern of Egr-1 expression in the atherosclerotic lesion, as well as the cell types involved, both carotid lesions and IMA were immunostained with an Egr-1–specific antibody. A total of 12 carotid lesions were examined as frozen sections (Figure 4). Staining in the lesion was strong in 9 of 12 patients with principal involvement of the α-actin–expressing SMC-like cells (Figure 4a). In contrast, the adjacent media, when present, showed much lower expression of Egr-1, though positive cells could be identified (Figure 4c). Regions of endothelial cells covering the lesion were occasionally positive for Egr-1. Double staining of macrophages with HAM56 and Egr-1 indicated that macrophages could be positive for Egr-1, but that Egr-1 staining was typically intense in the SMC-like cells surrounding the macrophages (not shown). Double-staining with α-actin, a SMC/myofibroblast marker, and Egr-1 indicated that the majority of Egr-1–positive cells were also α-actin positive (Figure 4b).

In contrast to the carotid lesions, the IMA showed much lower immunoreactivity. Five of 6 arteries showed no detectable staining for Egr-1 (Figure 4d), whereas 1 artery showed weak and diffuse staining in the medial SMC. Consistent with the RT-PCR findings, Egr-1 staining in the IMA was lower, or at best, similar to the weak, sparse staining of the carotid media shown in Figure 4c.

Figure 4
Immunocytochemical identification of Egr-1 in human atherosclerotic carotid arteries and IMA. Frozen sections of human endarterectomies or IMA were probed with Egr-1–specific antibodies. (a) Atherosclerotic carotid artery lesion stained with anti-Egr-1 antibodies yielding a dark brown reaction product in positive cells. (b) Double-staining reveals strong Egr-1 staining (brown) in lesion cells of the fibrous cap that co-stain with the SMC/myofibroblast marker α-actin (dark blue). Egr-1 staining is also visible in cells that stain only faintly for α-actin. (c) Adjacent tunica media from the same section showing sparse labeling. (d) Egr-1 staining in frozen sections from IMA obtained during bypass grafting. Only a light hematoxylin stain (blue) is visible, whereas Egr-1 staining (brown) is not detected.

Figure 5
The mRNA levels of Egr-1–responsive genes in the human lesion versus adjacent media. Human carotid artery lesions were dissected and mRNA was prepared as in Figure 1. The cDNA arrays were examined for the expression of Egr-1–inducible genes with the expression level expressed as a percent of housekeeping genes, as shown in Figure 1. Error bars, SEM; n = 3 experiments.
bers of the Egr-1 target gene family. Elevated levels of TNF-α (4-fold), IL-2 (16-fold), and M-CSF-1 (5-fold) transcripts were observed within the lesion relative to the media of the atherosclerotic vessel (Figure 5). PDGF-A chain was elevated 4-fold in the lesion from 3 to 12%. Both the α and β receptors for PDGF were also elevated by approximately 5-fold each (not shown), though their promoters are not known to be controlled by Egr-1. TGF-β1 and TGF-β3, both containing multiple predicted Egr-1 sites in their promoters (8, 21), were elevated in the lesion.

Egr-1–dependent genes may also include those whose products influence cell survival or provide protective functions. Levels of human copper-zinc superoxide dismutase (Cu, Zn-SOD) were increased in the lesions relative to the media about 5-fold (Figure 5). Analysis of the genomic arrays reveals that levels of p53 transcripts are increased in the lesion, relative to the media (Figure 5). In contrast, transcript levels of other transcription factors capable of regulating cell survival can be identified in end-stage, human carotid artery lesions. To examine the relationship of Egr-1 to the early development of lesions, the expression of the transcription factor in the vessel wall of homozygous LDLR−/− mice was examined immunocytochemically. The LDLR−/− mice fed a normal chow diet have only a modest elevation in plasma cholesterol and do not develop lesions (20). However, when fed a cholesterol-rich diet, these mice develop complex lesions of atherosclerosis similar to those in humans (23).

To determine the pattern of Egr-1 expression in the LDLR−/− model of atherosclerosis, mice were placed on the cholesterol-rich diet for 0, 5, 10, or 20 weeks. These time points were selected based on preliminary studies that demonstrated lesions begin to develop in the aortic arch and near the ostia after about 4 weeks on the diet. Sections of the aorta were immunostained with an Egr-1 antibody preblocked with an excess of peptide antigen (e) Aorta stained with Egr-1 antibody preblocked with irrelevant purified IgG.
at 10 and 20 weeks on the diet (Figure 6, c and d, respectively). In control studies, staining with the anti-Egr-1 antibody could be blocked by an excess of the Egr-1 peptide to which the antibody was raised (Figure 6e), and irrelevant purified IgG did not stain the sections (Figure 6f).

Expression of Egr-1 in atherosclerotic lesions formed at 20 weeks is shown in Figure 7. Egr-1 antigen can be detected in vascular SMC adjacent to the developing plaque (Figure 7a), as well as within cells found near the necrotic core (Figure 7b). However, little expression was detected in the macrophage-rich core. Thus, hypercholesterolemia-associated upregulation of Egr-1 expression suggests a role for this transcription factor in lesion formation.

Discussion
The lesions of atherosclerosis are thought to result from an excessive fibroproliferative response to multiple forms of insult to the vessel wall (19). To identify similarities and differences between the SMC-like cells of the lesion versus the “true” SMC of the arterial intima and media, these tissues were analyzed by cDNA expression arrays. Transcript profiling revealed that several genes were differentially regulated in the diseased vessels, most notably the transcription factor Egr-1. Multiple approaches were used to confirm that the induction of Egr-1 was valid: first, RT-PCR analysis of the same and similar tissues confirmed the findings from the arrays; second, elevated Egr-1 protein was detected by immunostaining in the atherosclerotic lesions; third, Egr-1 induction was observed in a mouse model of atherosclerosis; and finally, the expression of Egr-1–inducible genes was elevated in the same arrays, suggesting that Egr-1 induction led to transcription of Egr-1–responsive genes. All of these approaches suggest that the induction of Egr-1 and Egr-1–responsive genes in the human lesion is strong, reproducible, and potentially significant to the etiology of atherosclerosis.

There are certain inherent limitations to the expression analysis of RNAs from complex tissues like the atherosclerotic plaque. Foremost is that the lesion may contain multiple cell types that would differ from patient to patient. By focusing on the fibrous cap of the lesion, the present analysis would minimize, though not eliminate, the contribution of T cells and macrophages in the lesion because the majority of these cells tend to localize to “shoulder” regions and the fibro-fatty core of the plaque, both of which were intentionally trimmed at the same time that the endothelium was scraped.

The high level of Egr-1 in lesions could be viewed as surprising because the atherosclerotic plaque has a relatively low mitotic index, and the normal media has an even lower mitotic rate. In human carotid artery tissue acquired by endarterectomy, BrdU incorporation indicated that between 3 and 15% of the cells in the lesion might be proliferating, and many of these cells were identified as subendothelial macrophages (24). Considering that the immunocytochemistry indicated that the majority of the SMC-like cells were expressing elevated Egr-1, it would seem likely that the Egr-1 is expressed by nonproliferating cells that are in an activated state.

Egr-1 is readily induced by shear stress and laminar flow disturbances (25), mechanical injury, hypoxia, reactive oxygen species, and growth factors such as FGF and PDGF (3). Of particular relevance to atherosclerosis is the recent observation that both native and oxidized lipoproteins induce tissue factor gene expression in SMC through Egr-1 and Sp1 (26). Each of these stimuli might be relevant initiators of Egr-1 expression within the atherosclerotic lesion. Reactive oxygen species, in conjunction with released growth factors, might explain the locally high levels of Egr-1 expression in the SMC surrounding macrophage-rich regions of the plaque.

The induction of Egr-1 after endothelial injury in vitro appears to be due to local release of FGF-2 (17). In human atheroma, FGF-2 is readily expressed in the lesion (27), and infusions of FGF-2 markedly increase

Figure 7
Egr-1 expression in the late lesions of LDLR−/− mice fed a cholesterol-rich diet. LDLR−/− mice sacrificed at 20 weeks, were stained with Egr-1 antibodies. Regions containing raised atherosclerotic lesions are shown in (a) and (b) with Egr-1–positive cells producing a red reaction product (Fast Red) against the light-blue/green counterstain (methylene green).
intimal thickening after balloon injury of the rat (28). Likewise, NGF and its receptors are expressed in the human atherosclerotic lesion and are increased in response to vascular injury (29). Like PDGF, neurotropins have chemotactic actions on SMC and, in light of the current data, may augment expression of Egr-1 in the vascular lesion. Transcriptional control of Egr-1 expression appears to be positively regulated by a single, proximal serum response element (SRE) and a NF-κB–like negative regulatory element (30).

Egr-1 activity is probably closely regulated under normal conditions. Egr-1 can associate with corepressor proteins that can diminish the transcription of multiple target genes. These proteins include Egr-1/NGFI-A binding corepressors NAB1 and NAB2 (31–33). NAB2 is rapidly and transiently expressed in vascular SMC in response to mechanical injury or growth factors such as FGF-2 (34). These corepressors may be part of a negative feedback mechanism that serves to downregulate Egr-1–mediated gene transcription in injured SMC.

Egr-1 and its target genes have been correlated with the development of vascular lesions. In the present analysis, 13 genes with known or suspected Egr-1 responses were included within the cDNA expression arrays. All but 1 of these genes (NF-κB, p105/p50 subunit), followed an expression pattern that paralleled the expression of Egr-1. Whereas the present analysis is purely correlational, it demonstrates this correlation between Egr-1 and its target genes in human vascular disease. Stronger evidence of the relationship between Egr-1 and target gene induction has already been provided for many of the genes by using reporter genes and supershift binding assays.

Growth factors are one of several targets of Egr-1. In vitro studies have shown that Egr-1 binds to Egr-1 recognition sites within the promoter of both the PDGF-A (6) and the PDGF-B chain (4), resulting in induction of the mRNAs. In the present studies, PDGF-A was elevated almost 4-fold, consistent with previous reports demonstrating elevated PDGF-A in the lesion myofibroblasts (35). PDGF-B chain was not included in the arrays, though previous publications have indicated increased PDGF-B mRNA in the myofibroblasts of human carotid lesions compared to the adjacent media (35).

Another putative Egr-1–inducible growth factor gene is TGF-β1. Both TGF-β1 and TGF-β3 contain predicted Egr-1–binding sites in their promoters, and both mRNAs were elevated 3-fold in the fibrous cap of the lesion relative to the adjacent media. Previous studies have documented that both human (36) and animal (37) vascular lesions overexpress TGF-β1 mRNA and protein and that inhibition of TGF-β1 reduces the vascular fibroproliferative response to injury in rats (38). Active TGF-β1 and its 2 receptors are expressed focally in these late atherosclerotic lesions and particularly in the fibrous cap of the lesion (39). The Egr-1 induction of PDGFs and TGF-β would contribute to SMC/myofibroblast migration, proliferation, and extracellular matrix accumulation that are the hallmarks of advanced atherosclerosis.

The induction of other cytokines by Egr-1 would control the influx of inflammatory cells and their subsequent activation. Previous studies have identified functional Egr-1 sites in the promoters of the TNF-α (9) and IL-2 genes (40–42). M-CSF-1 is predicted to be an Egr-1 target gene on the basis of promoter sequence (43), but direct experimental verification is absent presently. The increase in TNF-α in the lesion may activate the endothelial and SMC and play a role in regulating the survival of cells in the lesion (44). The increased production of IL-2 and M-CSF-1 may prolong the survival of recruited lymphocytes and macrophages, respectively. ICAM-1 has been shown previously to be expressed at high levels in endothelial cells, macrophages, and SMCs of the human atherosclerotic lesions (45), as well as in rabbit and mouse vascular lesions (46). Egr-1–dependent induction of ICAM-1, in conjunction with cytokines and growth factors, may also play a role in the recruitment of cells to the lesions.

Other Egr-1–dependent genes include those whose products may further influence cell survival in the atherosclerotic lesion. One of these target genes is the tumor suppressor protein, p53, an essential regulatory molecule in cell proliferation and apoptosis. This tumor suppressor is an essential molecule in cell proliferation and apoptosis and has been postulated to play a part in the development of atherosclerosis. In the present studies, p53 mRNA was elevated almost 12-fold in the lesion, a finding supported by previous studies that observed elevated p53 immunoreactivity in human atherosclerotic lesions compared with IMA (47).

In the apoE-null mouse model of atherogenesis, the absence of p53 accelerates atherosclerosis by increasing cell proliferation in vivo, although p53-independent mechanisms appear to mediate apoptosis (48). Interestingly, in a human tumor cell line, Egr-1 counteracted apoptosis by the coordinated activation of TGFβ1, fibronectin, the cyclin-dependent kinase inhibitor, p21, and focal adhesion kinase, leading to enhanced cell attachment and reduced caspase activity (49). The survival effect of Egr-1 in this tumor line was dominant over the apoptotic effect of p53. Thus, Egr-1 may coordinate the expression gene products important for survival of vascular cells in atherosclerotic lesions.

Egr-1–dependent gene expression may also play a role in the control of oxidant stress in atherosclerotic lesions. SOD1 participates in the control of intracellular concentrations of reactive oxygen intermediates, and a functional Egr-1 element has been defined in the proximal promoter of the gene (50). Thus, Egr-1 responsiveness to reactive oxygen species leads to induction of SOD1, thereby completing a corrective feedback loop to control free-radical damage to the cell.

Egr-1 is commonly found to bind to promoter regions overlapping consensus Sp1 sites that can either facilitate or suppress transcription (51). In addition to the genes discussed above, known Egr-1–inducible genes include apoAI (52), 5-lipoxygenase (53), thymidine kinase (54), and the multidrug-resistance gene
MDR1 (55), although these genes were not contained within the arrays. Other genes, however, are thought to be suppressed by Egr-1, presumably by displacement of Sp1. For example, insulin-induced suppression of malic enzyme gene transcription appears to be mediated by Egr-1 (56). Likewise, B-cell clones overexpressing Egr-1 showed decreased expression of fas (CD95) and CD23, which makes them functionally resistant to fas-mediated apoptosis (57).

Given that Egr-1 is a stress-responsive gene, the present data indicates that the SMC/myofibroblast cells of the human and mouse lesions exist in a highly activated state, as opposed to passively accumulating in the intima. The elevations of Egr-1 in the human carotid lesion indicate that previous in vitro and in vivo studies implicating Egr-1 in the vascular response to injury can reasonably be extended to the human disease. Elevated Egr-1 expression in the lesion was associated with elevations in a wide spectrum of Egr-1–responsive genes that would be relevant to atherogenesis. As suggested by the recent in vivo studies of Santiago et al. (18), it is not unreasonable to speculate that interference with Egr-1 induction or function might prevent the negative sequelae of vascular injury after revascularization procedures such as angioplasty and endarterectomy.

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