The subendothelial extracellular matrix modulates NF-κB activation by flow: a potential role in atherosclerosis

A. Wayne Orr, John M. Sanders, Melissa Bevard, Elizabeth Coleman, Ian J. Sarembock, and Martin Alexander Schwartz

1Department of Microbiology and 2Department of Biomedical Engineering, 3Cardiovascular Research Center, 4Internal Medicine, and 5Mellon Prostate Cancer Research Center, University of Virginia, Charlottesville, VA 22908

Atherosclerotic plaque forms in regions of the vasculature exposed to disturbed flow. NF-κB activation by fluid flow, leading to expression of target genes such as E-selectin, ICAM-1, and VCAM-1, may regulate early monocyte recruitment and fatty streak formation. Flow-induced NF-κB activation is downstream of conformational activation of integrins, resulting in new integrin binding to the subendothelial extracellular matrix and signaling. Therefore, we examined the involvement of the extracellular matrix in this process. Whereas endothelial cells plated on fibronectin or fibrinogen activate NF-κB in response to flow, cells on collagen or laminin do not. In vivo, fibronectin and fibrinogen are deposited at atherosclerosis-prone sites before other signs of atherosclerosis. Ligation of integrin α2β1 on collagen prevents flow-induced NF-κB activation through a p38-dependent pathway that is activated locally at adhesion sites. Furthermore, altering the extracellular matrix to promote p38 activation in cells on fibronectin suppresses NF-κB activation, suggesting a novel therapeutic strategy for treating atherosclerosis.

Introduction

Atherosclerotic plaque develops in response to a localized inflammatory reaction in the vessel wall (Ross, 1999). Although known risk factors for atherosclerosis, such as hyperlipidemia, hypertension, diabetes, and smoking, play major roles in the incidence and progression of the disease, these factors are uniform throughout the circulation. The observation that atherosclerotic plaque forms preferentially at sites of disturbed blood flow (VanderLaan et al., 2004) suggests that flow patterns can regulate the chronic inflammation associated with atherogenesis (Caro et al., 1969; Ku et al., 1985; Glagov et al., 1988). In support of this idea, application of prolonged laminar flow to endothelial cells in culture is antiinflammatory and atheroprotective, whereas disturbed flow stimulates endothelial cell turnover, expression of prothrombotic and proinflammatory proteins, and altered redox regulation (Topper et al., 1996; Mohan et al., 1997; De Keulenaer et al., 1998; Brooks et al., 2002). These data suggest that disturbed flow patterns initiate the local inflammatory reaction seen during atherosclerosis, with amplification of lesions by humoral risk factors driving progression of the disease.

The NF-κB family of transcription factors are involved in numerous cellular processes, including differentiation, inflammation, proliferation, and apoptosis, and are postulated to contribute to atherogenesis (Collins and Cybulsky, 2001; Kutuk and Basaga, 2003). Current data implicate NF-κB as a key regulator of shear stress–induced inflammatory gene expression. NF-κB dimers, particularly the p50/p65 heterodimer, bind to a shear stress responsive element found in the promoter of several atherogenic genes, including ICAM-1, VCAM-1, and MCP-1 (monocyte chemotactic protein-1), which stimulate smooth muscle growth and migration (Resnick et al., 1993; Khachigian et al., 1995; Huo and Ley, 2001). NF-κB expression and activity, as well as ICAM-1 and VCAM-1 expression, are elevated in atherosclerosis-prone regions before or in the absence of fatty streak formation, indicating that they are very early events in atherosclerotic progression (Brand et al., 1996; Nakashima et al., 1998; Iiyama et al., 1999; Wilson et al., 2000). In vitro, disturbed shear stimulates prolonged NF-κB activation and NF-κB–dependent gene expression; in contrast, acute onset of

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Laminar shear stress activates NF-κB but only transiently (Lan et al., 1994; Khachigian et al., 1995; Mohan et al., 1997). We hypothesize that these differences in signaling pathways induced by laminar versus disturbed flow are due in part to differences in adaptation mechanisms, such that changes in flow velocity and direction associated with disturbed flow prevents down-regulation of the responses so that signals activated transiently by laminar flow are sustained in disturbed flow. Thus, data from both in vivo and in vitro models suggest that NF-κB contributes to the initiation of atherosclerosis by fluid shear stress.

Studies in vitro have shown that acute onset of laminar shear triggers conversion of integrins to a high affinity state, followed by their binding to the subendothelial ECM (Tzima et al., 2001). Resultant integrin signaling mediates activation of NF-κB through the small GTPase Rac (Tzima et al., 2002). Endothelial cells normally reside on a basement membrane comprised mainly of collagen (Coll) IV and laminin (LN). Coll binds primarily integrins α2β1 and α1β1, whereas LN binds mainly α6β4 (Belkin and Stepp, 2000; Heino, 2000). Ligation of these integrins is associated with a quiescent cell phenotype, consistent with the low turnover observed in endothelial cells in vivo (Schwartz and Assoian, 2001). Inflammation or injury can trigger the deposition of transitional ECM proteins such as fibronectin (FN) and fibrinogen (FG) into the subendothelial matrix (Sechler et al., 1998). In endothelial...
cells, FN primarily ligates α5β1 whereas FG ligates αvβ3, although other integrins may also be involved. Signals from α5β1 and αvβ3 are associated with enhanced endothelial cell proliferation and migration, processes important for injury-induced endothelial remodeling (Schwartz and Assoian, 2001). Many studies have shown that different integrins transduce distinct signals (Schwartz and Assoian, 2001). The involvement of integrins in shear stress–dependent signaling thus suggests that the composition of the subendothelial ECM may modulate cellular responses to fluid flow. Therefore, we investigated the role of the subendothelial ECM in the endothelial response to fluid shear stress.

**Results**

**Subendothelial matrix regulates shear stress–induced NF-κB activation**

To test whether or not shear stress–induced NF-κB activation depends on the underlying matrix, bovine aortic endothelial (BAE) cells plated on glass slides coated with Coll I, LN, FN, or FG for 4 h were subjected to laminar shear stress (12 dyne/cm²) in a parallel plate flow chamber. NF-κB is normally held inactive in the cytoplasm through interactions with IκB, but stimulus-induced IκB degradation results in NF-κB nuclear targeting and initiation of transcription (Chen and Greene, 2004; Yamamoto and Gaynor, 2004). As previously reported for endothelial cells under standard conditions (Scatena et al., 1998; Klein et al., 2002; Guo et al., 2004), cells on FN or FG showed enhanced nuclear translocation of NF-κB after flow; in contrast, cells on LN responded poorly and cells on Coll showed a higher baseline under static conditions, which decreased in response to flow (Fig. 1 A).

In addition to its nuclear targeting, numerous posttranslational modifications contribute to NF-κB’s transcriptional activity (Wang and Baldwin, 1998; Yang et al., 2003). Phosphorylation of the p65 subunit of NF-κB on Ser536 in the transactivation domain alters NF-κB–dependent transcription, presumably by regulating its interaction with other cis-acting elements. In addition, this site is phosphorylated by the IκB kinase (IKK) complex, and its phosphorylation is inhib-
ited by interaction with IκB, suggesting it is a good measure for pathway activation. Ser536 phosphorylation was stimulated in cells on FN or FG, but showed only small changes on LN; on Coll, NF-κB phosphorylation showed an elevated baseline that decreased after shear stress (Fig. 1B). Cells on Coll IV behaved identically to Coll I (unpublished data). This result was expected because Coll I and IV bind the same integrins (Defilippi et al., 1991; Tulla et al., 2001). To test if these matrix-specific signals translate to altered gene expression, we assayed expression of ICAM-1, an NF-κB–responsive gene. Both Western blotting and surface staining were performed in human umbilical vein endothelial cells due to poor antibody cross-reactivity with bovine ICAM-1. Both Western blotting and surface staining were performed in human umbilical vein endothelial cells due to poor antibody cross-reactivity with bovine ICAM-1. Both methods showed that, in cells on Coll, ICAM-1 expression diminished after shear stress, whereas expression increased in cells on FN and FG (Fig. 1C). Thus, NF-κB nuclear translocation, phosphorylation, and target gene expression correlate closely. We conclude that the transitional ECM proteins FN and FG support flow-induced NF-κB activation whereas LN and Coll do not.

**ECM remodeling in vivo**

The distinct effects of different ECM proteins in flow-induced NF-κB activation in vitro prompted us to ask whether or not these effects were relevant to plaque development in vivo. To test whether FN and FG deposition are associated with focal expression of NF-κB target genes in atherosclerosis-prone regions of the vasculature, these proteins were examined in mice. Serial sections from atherosclerosis-susceptible sites were stained for ECM proteins, for ICAM-1 and VCAM-1 as markers of NF-κB activation, and for the leukocyte integrin Mac-2 as a measure of monocyte recruitment. Atherosclerosis-resistant C57/B6 mice and atherosclerosis-prone apolipoprotein E (ApoE) null mice were fed either a normal chow diet or a high fat, Western diet for 10 wk. Sites associated with atherosclerotic plaque formation, such as the innominate artery and the carotid arteries, were then analyzed by immunohistochemistry (IHC; Nakashima et al., 1994).

In C57/B6 mice fed the low fat, chow diet, sites that are susceptible to atherosclerosis in other models demonstrated focal increases in ICAM-1 and VCAM-1 staining and in-
increased staining for FN, although no Mac-2 staining was apparent (Fig. 2A). We did not detect any change in FG (Fig. 2A) or LN (not depicted) staining and no staining for FN in regions outside these zones. ApoE null mice fed a chow diet have elevated cholesterol and develop atherosclerotic lesions over the course of their lifespan (Daugherty, 2002). In these mice, similar colocalization of ICAM-1, VCAM-1, FN, and FG was observed when nearby sections were stained (Fig. 2B). FN has been reported in well-defined fatty streaks (Stenman et al., 1980; Labat-Robert et al., 1985; Shekhoon et al., 1987; Tanouchi et al., 1992), but these data are the first evidence that FN is deposited before fatty streak development. In contrast, no changes in PECAM staining were evident (unpublished data). VCAM-1 staining in vascular smooth muscle cells is indicative of smooth muscle activation and is associated with atherosclerosis, which is consistent with our interpretation of these regions as early atherogenesis (Li et al., 1993).

ApoE null mice fed a Western diet develop atherosclerosis at greatly accelerated rates (Daugherty, 2002). Under these conditions, FN, FG, VCAM-1, and ICAM-1 staining was strongly enhanced in atherosclerosis-prone sites compared with C57/B6 or ApoE null mice fed a chow diet; however, no Mac-2 staining was apparent (Fig. 2C). As before, these changes were restricted to atherosclerosis-prone regions near bifurcations. Together, these data show that ECM is remodeled in regions of disturbed flow in vivo before other indicators of atherosclerosis and that NF-κB activation correlates with these changes in the subendothelial ECM. Furthermore, conditions favoring hypercholesterolemia, such as depletion of ApoE or a high fat diet, enhance matrix remodeling specifically at sites of disturbed flow, suggesting a potential synergy between the atherogenic effects of hypercholesterolemia and disturbed flow in vivo.

Coll-specific p38 signaling inhibits shear-induced NF-κB activation

In addition to NF-κB, p38 MAP kinase is also involved in stimulating inflammatory protein expression in a variety of systems. However, p38 shows differential effects on NF-κB, either enhancing or suppressing NF-κB depending on the stimulus (Schwenger et al., 1998; Alpert et al., 1999; Bowie and O’Neill, 2000; Bradbury et al., 2001). To examine p38 in this system, BAE cells were plated on different ECM proteins for 4 h and effects of shear were tested. Western blotting with an antibody against the phosphorylated and activated form of p38 showed that shear stress stimulated a transient increase in p38 phosphorylation in cells on Coll but not on other ECM proteins (Fig. 3A). This result is consistent with previous reports showing that the Coll-binding integrin α2β1 stimulates p38 activation (Ivaska et al., 1999). To test if this Coll-specific p38 activation mediates inhibition of shear stress–induced NF-κB activation, cells were preincubated with the p38 inhibitor SB202190 restored shear stress–induced NF-κB activation, cells were preincubated with the p38 inhibitor SB202190 (1 μM for 1 h) and the phosphorylation of p65 was assessed by immunoblotting. Bands were quantified by densitometry and normalized for total p65 (Fig. 3B, C). Cells on Coll I or FN were pretreated with the p38 inhibitor SB202190 (1 μM for 1 h) and integrin ligation was induced with the β1 integrin–activating antibody TS2/16 (15 μg/ml for 1 h). p65 localization was assessed by immunostaining, and the percentage of cells showing nuclear localization was scored (n = 3; *, P < 0.05). (D) TS2/16-induced p65 phosphorylation of p65 on Ser536 was assessed by Western blotting. Bands were quantified by densitometry and normalized to total p65 (n = 3–6).

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blocking α2β1 antibody 12F1 (20 μg/ml for 30 min). With this short exposure to blocking antibodies, we can inhibit new integrin binding without significantly affecting existing adhesions or cytoskeletal structure (Fig. S1, available at http://www.jcb.org/cgi/content/full/jcb.200410073/DC1). Blocking α2β1 integrin ligation prevented the shear stress–induced activation of p38 in cells on Coll (Fig. 4 A) and rescued NF-κB phosphorylation (Fig. 4 B). The control antibody had no effect. Preventing new ligation of integrin α6β1 with the blocking antibody GoH3 (20 μg/ml for 30 min) did not enhance flow-induced p65 nuclear translocation or phosphorylation in cells on LN, instead, the already low level of NF-κB was reduced further. These results suggest α6β1 is not suppressive (Fig. S2, A–C, available at http://www.jcb.org/cgi/content/full/jcb.200410073/DC1).

To test whether or not activating integrins, as occurs in response to fluid flow, is sufficient to induce these pathways, cells were treated with the β1 integrin–activating antibody TS2/16 (15 μg/ml). Antibody-induced integrin activation resulted in a similar, matrix-specific nuclear translocation (Fig. 4 C) and phosphorylation (Fig. 4 D) of NF-κB on FN but not Coll. Furthermore, TS2/16-induced NF-κB activation on Coll was rescued by the p38 inhibitor SB202190 (Fig. 4, C and D). These data support the conclusion that integrin activation and ECM binding initiate these pathways.

**Flow-induced NF-κB activation on mixed matrices**

Cells in vivo are usually exposed to multicomponent matrices of varying composition. To test if Coll can show dominant effects in this system, cells were plated on increasing concentrations of FN in either the absence or presence of an underlying Coll matrix. The amount of FN adsorbed to the coverslips was assayed in separate experiments. In the absence of Coll, p65 nuclear translocation and Ser536 phosphorylation displayed a dose-dependent increase as FN concentration increased (Fig. 5, A and B). The dose-dependent effect reached a maximum between 4 and 6 μg of deposited FN, which corresponds to 20 μg/ml in the coating solution. When examined on coverslips first coated with Coll, the amount of FN necessary to elicit NF-κB activation was notably higher, and maximal p65 activation was reduced even at the highest doses of FN. These results support the notion that Coll signaling actively suppresses the FN-dependent activation of NF-κB by flow. However, suppression of signaling may work in the opposite direction because FN deposition inhibited flow-induced p38 phosphorylation in cells on Coll in a dose-dependent manner (Fig. 5 C).

**Signaling is localized to sites of adhesion**

To test if elevated p38 activity on Coll suppresses NF-κB activation in a more general way, cells were treated with TNFα, a cytokine that strongly induces this pathway. Interestingly, no suppression of TNFα-induced NF-κB was observed (Fig. S3 A, available at http://www.jcb.org/cgi/content/full/jcb.200410073/DC1). Furthermore, transfection with a constitutively active MKK3 (MKK3 EE), a kinase that is upstream of p38 and activates p38 globally (not depicted), did not prevent the shear stress– or TNFα-induced activation of NF-κB in cells on FN (Fig. S3 B).

These results appear paradoxical but could be explained if the inhibitory p38 signal from Coll occurs locally within a specific compartment. To test this hypothesis, endothelial cells on Coll or FN were sheared for 5 min and active p38 was localized. Cells on Coll showed colocalization of phospho-p38 with β1 integrins, which was increased by shear stress (Fig. 6 A). Cells on FN showed only low levels of phospho-p38 staining, which is consistent with results from Western blotting.

The IKK complex consists of two kinase subunits (IKKα and IKKβ) and a regulatory subunit named NEMO or IKKγ (Israel, 2000). Numerous stimuli activate NF-κB through this complex, resulting in phosphorylation of the kinase subunits (Ser176/
Ser180 for IKKα or Ser177/Ser181 for IKKβ (Israel, 2000). Shear stress stimulates NF-κB activation through the integrin- and Rac-dependent activation of IKKα/β, possibly through Nck-interacting kinase (Bhullar et al., 1998; Tzima et al., 2002; Hay et al., 2003). Immunoblotting with phospho-specific antibodies reveals that shear stress stimulates a matrix-specific phosphorylation profile of IKKβ (Fig. 6 B) that correlated well with activation of NF-κB. These results suggest that the divergent matrix-specific signaling occurs upstream of IKK activation. Curiously, shear stress–induced IKK phosphorylation appeared to specifically target IKKβ. This result may be related to recent findings that IKKβ is the primary component involved in NF-κB activation, whereas IKKα has distinct functions (Karin et al., 2004). Staining of fixed cells on Coll with antibody to phospho-IKK showed distinct colocalization of phospho-IKK with β1 integrins in unsheared cells, which was diminished after exposure to shear stress (Fig. 6 C). In cells on FN, phospho-IKK also colocalized with β1 integrins, and this signal increased after shear stress. In contrast, TNFα-induced IKK phosphorylation resulted in diffuse cytoplasmic and nuclear staining (Fig. 6 D). These results suggest that the localized activation of p38 within integrin adhesions specifically inhibits the local, but not global, activation of NF-κB.

### Altering the FN matrix to inhibit NF-κB

The selective activation of NF-κB in atherosclerosis-prone regions of the vasculature, together with the importance of NF-κB target genes in atherosclerosis, suggest that inhibition of this transcription factor could have therapeutic value. However, NF-κB has important roles in cell survival and immune protection, thus, global inhibition of the pathway is likely to be deleterious. In contrast, local inactivation of NF-κB within cell–ECM adhesions in atherosclerotic plaque could be beneficial. Although FN does not normally activate p38, a peptide derived from the first type III repeat in FN, termed III-1C, alters the

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**Figure 6. Localized activation of p38 and IKK at adhesion sites.**

(A) Cells were plated on Coll or FN, sheared for 5 min or kept under static conditions, and stained for phosphorylated p38 and β1 integrin. Images are representative of four experiments. (B) BAE cells were plated on Coll, LN, FN, or FG and sheared for the indicated times, and IKK phosphorylation was assessed by Western blotting. Results are representative of four to six experiments. (C) Cells were plated on Coll or FN, sheared for 30 min or kept as static controls, and stained for phosphorylated IKK and β1 integrin. Results are representative of three experiments. (D) Cells on Coll or FN were treated with 10 U/ml TNFα and stained for phosphorylated IKK and β1 integrin. Images are representative of three experiments.
structure of the FN matrix and stimulates p38 activation (Yi and Ruoslahti, 2001; Klein et al., 2003). When cells on FN were treated with this peptide, activated p38 colocalized with integrins at sites of adhesion (Fig. 7, A and B). To test whether or not FNIII-1C might also suppress NF-κB activation, BAE cells on FN were treated for 4 h with FNIII-1C or as controls with heat denatured FNIII-1C or the analogous peptide from the second type III repeat, FNIII-2C. Cells were sheared for 30 min, and NF-κB nuclear translocation was assessed by immunocytochemistry (n = 3–5). (D) Cells were pretreated with FN peptides as in C, followed by incubation with the p38 inhibitor SB202190 (1 μM) for 1 h before onset of shear stress. Cells were sheared for 30 min and nuclear translocation of p65 was assessed. Values are means ± SD (>100 cells counted per condition per experiment; n = 3–8), **, P < 0.01; ***, P < 0.001. (E) Cells were pretreated with the FN peptides as indicated, incubated with SB202190, and sheared for 30 min. p65 phosphorylation at Ser536 was assessed by Western blotting. Bands were quantified by densitometry and normalized to total p65. Values are means ± SD from four to seven experiments. *, P < 0.05; **, P < 0.01; ***, P < 0.001.

Discussion

We present evidence that ECM composition leading to specific integrin signals can determine the activation of NF-κB by shear stress. We also show that comparable ECM remodeling occurs in vivo and correlates with activation of NF-κB in regions of...
disturbed shear stress. Therefore, these results suggest that this interplay might control early atherogenesis in vivo.

The role of the extracellular matrix as a key regulator of the behavior of advanced atherosclerosis is widely accepted, with ECM remodeling thought to affect both the size and stability of the fibrous atherosclerotic plaque (Katsuda and Kaji, 2003). However, a critical role for the subendothelial matrix in development of atherosclerosis has not been suspected. Our data show that deposition of FN and FG into the subendothelial matrix in regions of disturbed flow precedes fatty streak formation in vivo. These proteins in the subendothelial matrix promote activation of NF-κB by shear stress in vitro, whereas the normal basement membrane component Coll suppresses NF-κB through α2β1-dependent p38 activation. Importantly, p38 is activated locally within adhesion sites and prevents the shear stress–induced activation of IKKβ and NF-κB but not the global activation of NF-κB by cytokines. Finally, we demonstrate that FN matrix can be altered to stimulate p38 and suppress shear stress–induced NF-κB activation, suggesting that FN is a possible target for therapeutic intervention.

**ECM remodeling in atherogenesis**

The normal subendothelial ECM is a basement membrane whose major integrin-binding components are type IV Coll and LNs (Yurchenco and O’Rear, 1994). Injury, inflammation, or angiogenesis result in degradation of the basement membrane and deposition of transitional ECM proteins such as FN and FG. We show here that the presence of transitional matrix proteins in the subendothelial matrix correlates with NF-κB activation and target gene expression in regions of disturbed shear stress. These data are consistent with the idea that transitional matrix deposition regulates inflammatory gene expression in early atherogenesis; however, the causal relationships in vivo between ECM and NF-κB remain to be elucidated. Effects of disturbed flow on FN matrix deposition could involve changes in endothelial permeability to plasma proteins (Friedman and Fry, 1993; Phelps and DePaola, 2000; Himburg et al., 2004), changes in endothelial expression of ECM genes (Brooks et al., 2002), or changes in integrin expression or function (Brooks et al., 2002). Identifying the mechanisms involved in local matrix remodeling at atherosclerosis-prone sites in arteries will be an interesting direction for future work.

We also observed marked increases in both the ECM proteins FN and FG and the inflammatory markers ICAM-1 and VCAM-1 in atherosclerosis-susceptible regions of arteries from ApoE null mice fed a Western diet compared with mice on a Chow diet, suggesting that the Western diet may further increase ECM remodeling. This effect would be predicted to further enhance endothelial responses to disturbed flow, thus defining a positive feedback loop that would contribute to disease progression.

**Inhibition of NF-κB activation by locally activated p38**

Although p38 enhances NF-κB activity in many systems (Carter et al., 1999; Korus et al., 2002), there are several stimuli for which activation of p38 inhibits NF-κB, including vitamin C, salicylate, indomethacin, sorbitol, hydrogen peroxide, and arsenite (Schwenger et al., 1998; Alpert et al., 1999; Bowie and O’Neill, 2000; Bradbury et al., 2001). *Drosophila melanogaster* p38 isoforms have been shown to limit antimicrobial peptide production, a known target of *D. melanogaster* NF-κB (Han et al., 1998). In none of these cases has the mechanism by which active p38 represses NF-κB activation been elucidated. Binding of integrin α2β1 to Coll also activates p38 (Ivaska et al., 1999). Consistent with our model in which shear stress stimulates integrin activation and ligand binding (Jalali et al., 2001; Tzima et al., 2001; Katsumi et al., 2004), we found that shear stress stimulated α2β1-dependent p38 activation in cells on Coll. Activation of p38 inhibited both IKKβ and NF-κB activation. Importantly, both active p38 and active IKKβ localized to sites of adhesion. This colocalization correlated with the integrin specificity of the functional interaction because Coll did not inhibit cytokine activation of NF-κB and global activation of 38 did not inhibit shear stress activation of NF-κB. Therefore, the data provide a striking example where signaling specificity is conferred by compartmentalization. Future work will be required to elucidate the pathway by which p38 inhibits IKKβ in this system.

**Altering the FN matrix**

Selective inhibition of NF-κB activation by shear stress offers potential benefits for treatment of atherosclerosis, but it is essential that inhibitors act locally both to be effective and to avoid global inhibition of NF-κB, which would likely have undesirable side effects. A peptide from the first type III repeat of FN that was reported to trigger p38 activation in cells on FN induced local activation of p38 at sites of adhesion on a FN matrix and inhibited shear stress–induced activation of NF-κB. The effects of FNIII-1C are poorly understood. FNIII-1C has been reported to trigger assembly of soluble FN into fibrils and increase matrix FN (Morla and Ruoslahti, 1992), inhibit deposition of matrix FN (Bourdoulous et al., 1998), and alter its conformation without changing the total amount of matrix (Klein et al., 2003). Changes in FN matrix structure could conceivably alter signaling through a previously bound receptor, perhaps by changing its spatial organization, or could expose new cell binding sites or block existing cell binding sites required to prevent p38 activation. FNIII-1C binding to FN masks the EDA domain present in certain forms of FN (Klein et al., 2003), which is consistent with this last hypothesis. Curiously, mice engineered to lack the EDA domain of FN show reduced atherosclerosis, suggesting a potential role for this interaction in atherogenesis (Tan et al., 2004).

FNIII-1C itself is not a likely candidate for therapy, at least as an injected reagent, because it binds to soluble FN in the plasma and has been reported to induce assembly of this FN into fibrils, an event that could trigger thromboses or other deleterious events. Thus, further investigation into the mechanisms of ECM remodeling and p38 activation will be needed to develop more suitable methods for altering subendothelial ECM or cellular responses to it. Nevertheless, these results point toward an entirely novel strategy for treating atherosclerosis. The substrate dependence for proatherosclerotic gene expression may also have important implications in determining...
suitable signaling-based therapeutic approaches and designing bioengineered vascular grafts.

We conclude that transitional matrix proteins, such as FN and FG, are required for the flow-induced activation of NF-κB and suggest that deposition of FN and FG into the subendothelial matrix at atherosclerosis-prone sites in vivo may regulate the early changes in inflammatory gene expression associated with atherosclerosis. Furthermore, this shear stress–induced inflammatory signal appears to be inhibited by the endogenous Coll basement membrane through integrin α2β1–dependent activation of p38. Exploiting this fact, we show that alteration of the FN matrix with the FNIII-1C peptide prevents the shear stress–induced activation of NF-κB through localized p38 signaling. These results suggest a model in which subendothelial matrix remodeling mediates atherosclerotic progression through integrin-dependent mechanotransduction. Further work in which the subendothelial matrix or integrin signals are altered will be required to rigorously test this new model.

Materials and methods

Cell culture, parallel plate flow chamber, and transfection

BAE cells were maintained in DMEM containing 10% FBS, 1% penicillin/streptomycin, and 2 mM L-glutamin (Invitrogen). Cells were plated on 38 × 75 mm glass slides (Corning) precoated with 40 μg/ml Coll I, 20 μg/ml LN, 20 μg/ml FN, or 20 μg/ml FG. For mixed matrix experiments, slides coated with or without Coll I were subsequently coated with increasing concentrations of FN. Western blotting was used to determine the amount of FN adsorbed to the dishes with and without Coll coating. After 4 h, cells were fully attached and spread and formed a confluent monolayer. Slides were loaded onto a parallel plate flow chamber in either 0.1% BSA or 0.5% FBS and 12 dynes/cm² of shear stress was applied for varying times. Transient transfection of dominant-negative p38 (AGF mutation; 10 μg/ml) and constitutively active MKK3 (EE mutation; 10 μg/ml) was performed by electroporation at 170 V and 960 μF. Cells were replated and incubated for 48 h to allow protein expression before loading onto glass slides for exposure to shear stress.

Immunocytochemistry

Cells were fixed with PBS containing 2% formaldehyde, permeabilized with 0.2% Triton X-100 when applicable, and blocked overnight in PBS containing 1% BSA and 10% goat serum. Primary antibodies were incubated with cells in blocking buffer as follows: rabbit anti-p65 (Santa Cruz Biotechnology, Inc.; 1:250 for 1 h), rabbit anti-ICAM (Santa Cruz Biotechnology, Inc.; 1:100 for 1 h), rabbit anti–phospho-β1 integrins (1G210; 1 μg/ml for 1 h). Cells were incubated in 1 μg/ml Alexa 488–conjugated anti-rabbit IgG or Alexa 568–conjugated anti–mouse IgG (Molecular Probes). Slides were mounted with Fluoromount G and imaged with a Nikon TE300 microscope equipped with a CoolSnap video camera (Photometrics) using the Inovision ISEE software program.

Immunoblotting

Cell lysis and immunoblotting were performed as previously described (On et al., 2002). Rabbit anti–phospho-p38 (Cell Signaling Technology), mouse anti-p38 (Santa Cruz Biotechnology, Inc.), rabbit anti–phospho-p65 (Ser536; Cell Signaling Technology), rabbit anti-p65 (Santa Cruz Biotechnology, Inc.), rabbit anti–phospho-IKKα/β (Cell Signaling Technology), and rabbit anti–IKKα/β (Santa Cruz Biotechnology, Inc.) were all used at 1:1000 dilutions.

Animals

Six male ApoE-deficient mice on a C57BL/6 background (Jackson ImmunoResearch Laboratories), 8–12 wk old and weighing 18–20 g, were used in these experiments. Four mice were fed a Western-type atherogenic diet (TD 88137; Harlan-Teklad; containing 21% fat by weight, 0.15% by weight cholesterol, and 19.5% by weight casein without so-

dium cholate) for 10 wk before killing. Control mice were fed a Chow diet during this time. Comparable wild-type C57/B6 mice were fed the Chow diet during the same time frame.

Vessel harvest

At 20 wk old (10 wk on diet), mice were perfused with 4% formaldehyde and the innominate, left carotid, right carotid, and short segments of the descending abdominal aorta near the renal arteries and the iliac bifurcation were processed for paraffin embedding.

IHC

5-μm paraffin sections were obtained for IHC. IHC for adhesion molecules VCAM-1, ICAM-1, Mac-2, and PECAM-1 (Santa Cruz Biotechnology, Inc.) was performed as previously described (McPherson et al., 2001). After microwave antigen retrieval with antigen unmasking solution (Vector Laboratories), rabbit anti-FN (1:400; Sigma-Aldrich) and rabbit anti-LN (1:500; Sigma-Aldrich) were applied. Detection of antibodies was done with Vectastain Elite Kit (Vector Laboratories). Visualization was done with DAB (Deko Corp). Goat anti-FG (1 μg/ml; Accurate Chemical) was applied as for the other antigens, except that no antigen retrieval was required. Images were acquired using the 20 or 40× objective on a microscope (model BX51; Olympus) equipped with a digital camera (model DP70; Olympus) using the ImagePro Plus software program in the Academic Computing Health Sciences Center at the University of Virginia.

Online supplemental material

In Fig. 4, we demonstrate that incubation with the α2β1 integrin–blocking antibody R2-8C8 can block shear-induced p38 activation on Coll and rescue NF-κB activation in cells on Coll, presumably by preventing the interaction of newly activated α2β1 with Coll. To demonstrate that this effect was not due to changes in cell adhesiveness, we show that short-term incubation with R2-8C8 does not affect the organization of the actin cytoskeleton (Fig. S1). To show that this effect was specific to Coll, we also prevented the interaction between α2β1 and LN with the α6β1-blocking antibody GoH3, which did not rescue flow-induced NF-κB activation (Fig. S2). Finally, we show that flow-induced p38 activation on Coll prevents flow-dependent activation of NF-κB. To determine if this effect was specific for flow, the ability of TNFα to activate NF-κB in cells on different matrix proteins or overexpressing a constitutively active MKK3 construct (MKK3-EE) was assessed (Fig. S3). Online supplemental material is available at http://www.jcb.org/cgi/content/full/jcb.200410073/DC1.

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