Role of Endothelial Dysfunction in Atherosclerosis

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Abstract—As the major regulator of vascular homeostasis, the endothelium exerts a number of vasoprotective effects, such as vasodilation, suppression of smooth muscle cell growth, and inhibition of inflammatory responses. Many of these effects are largely mediated by nitric oxide, the most potent endogenous vasodilator. Nitric oxide opposes the effects of endothelium-derived vasoconstrictors and inhibits oxidation of low-density lipoprotein. A defect in the production or activity of nitric oxide leads to endothelial dysfunction, signaled by impaired endothelium-dependent vasodilation. Accumulating evidence suggests that endothelial dysfunction is an early marker for atherosclerosis and can be detected before structural changes to the vessel wall are apparent on angiography or ultrasound. Many of the risk factors that predispose to atherosclerosis can also cause endothelial dysfunction, and the presence of multiple risk factors has been found to predict endothelial dysfunction. A number of clinical trials have shown that 3-hydroxy-3-methylglutaryl coenzyme A (HMG-CoA) reductase inhibitors (statins) improve endothelial dysfunction in patients with coronary risk factors beyond what could be attributed to their impact on plasma lipids. Studies have elucidated several possible mechanisms by which statin therapy may improve endothelial dysfunction, including upregulation of nitric oxide production or activity and reduction of oxidative stress. (Circulation. 2004;109[supp III]:III-27–III-32.)

Key Words: atherosclerosis □ endothelial dysfunction □ nitric oxide □ statins

As the major regulator of vascular homeostasis, the endothelium maintains the balance between vasodilation and vasoconstriction, inhibition and stimulation of smooth muscle cell proliferation and migration, and thrombogenesis and fibrinolysis.1,2 When this balance is upset, endothelial dysfunction occurs, causing damage to the arterial wall. Endothelial dysfunction is considered an early marker for atherosclerosis, preceding angiographic or ultrasonic evidence of atherosclerotic plaque.1 The integral role of the endothelium in vascular health and of endothelial dysfunction in atherosclerosis has generated considerable interest in the potential for reversal of endothelial dysfunction with lipid-lowering therapy.

Regulatory Functions of the Endothelium

The normal, healthy endothelium regulates vascular tone and structure and exerts anticoagulant, antiplatelet, and fibrinolytic properties. The maintenance of vascular tone is accomplished by the release of numerous dilator and constrictor substances. A major vasodilative substance released by the endothelium is nitric oxide (NO), originally identified as endothelium-derived relaxing factor (EDRF). Other endothelium-derived vasodilators include prostacyclin and bradykinin.3 Prostacyclin acts synergistically with NO to inhibit platelet aggregation.1 Bradykinin stimulates release of NO, prostacyclin, and endothelium-derived hyperpolarizing factor, another vasodilator, which contributes to inhibition of platelet aggregation.3 Bradykinin also stimulates production of tissue plasminogen activator (t-PA), and thus may play an important role in fibrinolysis.

The endothelium also produces vasoconstrictor substances, such as endothelin (the most potent endogenous vasoconstrictor identified to date) and angiotensin II. Angiotensin II not only acts as a vasoconstrictor but is also pro-oxidant4 and stimulates production of endodin. Endothelin and angiotensin II promote proliferation of smooth muscle cells and thereby contribute to the formation of plaque.5 Activated macrophages and vascular smooth muscle cells, characteristic cellular components of atherosclerotic plaque, produce large amounts of endothelin.5

Damage to the endothelium upsets the balance between vasoconstriction and vasodilation and initiates a number of events/processes that promote or exacerbate atherosclerosis; these include increased endothelial permeability, platelet aggregation, leukocyte adhesion, and generation of cytokines.6 Decreased production or activity of NO, manifested as impaired vasodilation, may be one of the earliest signs of atherosclerosis.

Nitric Oxide

Nitric oxide is a pivotal endothelium-derived substance. The hallmark of endothelial dysfunction is impaired endothelium-dependent vasodilation, which is mediated by NO. A defect in NO production or activity has been proposed as a major predictor of atherosclerotic plaque formation.
mechanism of endothelial dysfunction and a contributor to atherosclerosis.

Production of Nitric Oxide
Nitric oxide is formed in endothelial cells from its precursor L-arginine via the enzymatic action of endothelial nitric oxide synthase (eNOS), which is located in caveolae (invaginations in cell membranes). The protein caveolin-1 binds to calmodulin to inhibit activity of eNOS; the binding of calcium to calmodulin displaces caveolin-1, activating eNOS and leading to production of NO. Cofactors such as tetrahydrobiopterin and nicotinamide adenine dinucleotide phosphate (NADPH) are also involved in NO production (Figure 1). The molecular basis of eNOS signaling is shown in greater detail in Figure 2.

Shear stress increases the expression of eNOS. Asymmetric dimethylarginine (ADMA) inhibits NO, and elevated levels of ADMA have been associated with endothelial dysfunction and atherosclerosis. The isoprenoid geranylgeranyl pyrophosphate, an intermediate factor in the cholesterol synthesis pathway, also inhibits synthesis of eNOS. Plasma and macrophage content of oxidized LDL increases synthesis of caveolin-1, which inhibits production of NO by inactivating eNOS. Oxidized LDL cholesterol increases synthesis of caveolin-1, which inhibits production of NO by inactivating eNOS.

Functions of Nitric Oxide
Nitric oxide mediates endothelium-dependent vasodilation by opposing the effects of endothelium-derived vasoconstrictors such as angiotensin II and endothelin. It also inhibits platelet adherence and aggregation, leukocyte adhesion/infiltration, and proliferation of vascular smooth muscle cells. Nitric oxide prevents oxidative modification of low-density lipoprotein (LDL) cholesterol. Oxidation of LDL has been proposed as a major mechanism of the atherosclerotic process; furthermore, plasma and macrophage content of oxidized LDL in coronary plaques correlate with severity of acute coronary syndrome. Conversely, impaired production or activity of NO leads to events or actions that promote atherosclerosis, such as vasoconstriction, platelet aggregation, smooth muscle cell proliferation and migration, leukocyte adhesion, and oxidative stress. Oxidized LDL cholesterol increases synthesis of caveolin-1, which inhibits production of NO by inactivating eNOS. Oxidative stress can also interfere with the production and activity of NO by a number of mechanisms that are independent of LDL. For example, the free radical superoxide anion rapidly inactivates NO and destroys tetrahydrobiopterin, a cofactor required for NO synthesis.

Clinical Assessment of Endothelial Function
Endothelial function can be assessed invasively using acetylcholine, which induces endothelium-dependent dilation and
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A study by Ludmer et al using the acetylcholine test provided the first evidence in humans of impaired endothelium-dependent vasodilation in the presence of atherosclerosis. Reactive hyperemia induces increased blood flow and shear stress, stimulating NO release and flow-mediated dilation (FMD) that can be quantified as an index of vasomotor function. The systemic nature of atherosclerosis is reflected by the close correlation between endothelial dysfunction in the forearm and coronary endothelial dysfunction. These findings also suggest that noninvasive assessment of peripheral arteries may be useful for determining the effects of risk factors on endothelial function and for evaluating the effects of therapy. Finally, endothelial function correlates inversely with serum C-reactive protein (CRP).

Endothelial Dysfunction Predicts Clinical Outcome

Research linking endothelial dysfunction with risk factors for coronary disease is intriguing and opens potentially useful avenues of investigation into the pathophysiology of coronary disease and also into the development of therapies targeting the endothelium. Given the links between endothelial dysfunction and coronary risk factors, it is not surprising that endothelial dysfunction is also associated with clinical events related to atherosclerosis.

A study in patients with mild (nonobstructive) CAD found that severe coronary endothelial dysfunction significantly increased the risk of cardiac events over an average follow-up of 28 months. In contrast, patients with mild dysfunction or normal endothelial function experienced no cardiac events. Evidence from subsequent studies reinforced the concept that coronary endothelial function may be a useful prognostic...
indicator. Coronary endothelial vasodilator dysfunction was found to independently predict progression of atherosclerosis and risk of cardiovascular events over a median follow-up of 7.7 years, even after the data were adjusted for conventional coronary risk factors.29 In the largest such study to date, epicardial and microvascular coronary endothelial dysfunction independently predicted acute cardiovascular events in patients with and without CAD.30

In a study in which vasodilation was evaluated using plethysmography of forearm blood flow in response to acetylcholine (endothelium-dependent) and sodium nitroprusside (endothelium-independent), patients who experienced cardiovascular events over a mean follow-up of 4.5 years showed impaired vasodilator responses.31 Another study32 monitored 73 patients who underwent cardiac catheterization for chest pain and evaluation of brachial artery FMD by high-resolution ultrasound over a mean of 5 years. Cardiovascular events including percutaneous and surgical revascularization occurred more often in patients with impaired FMD (<10%) compared with patients with preserved FMD (>10%).

Correction of Endothelial Dysfunction
A number of interventions have been shown to be effective in restoring endothelium-dependent vasodilation. These include lipid-lowering therapy (eg, 3-hydroxy-3-methylglutaryl coenzyme A [HMG-CoA] reductase inhibitors [statins], cholesterol, or LDL apheresis), angiotensin-converting enzyme inhibitors, antioxidants, reducing hyperglycemia, diet, and exercise. The effectiveness of dietary L-arginine on endothelial dysfunction in humans is not fully established and further research is needed.33 The following discussion focuses primarily on the role of statins in improving endothelial dysfunction.

Evidence for Beneficial Effects of Statins
As early as 1994, statin therapy was shown to improve endothelium-dependent dilation of coronary and peripheral arteries in patients with hypercholesterolemia. In one study, 6 months of treatment with pravastatin (10 to 20 mg/d) improved endothelium-dependent coronary vasomotion in patients with hypercholesterolemia.34 In another study of patients with atherosclerosis, cholesterol lowering with lovastatin (40 mg twice daily) significantly improved endothelium-mediated responses in the coronary arteries after 6 months.35

However, improvement has been detected even earlier after initiation of therapy. For example, in hypercholesterolemic, postmenopausal women who received atorvastatin (10 mg/d) for 8 weeks, a significant increase in endothelium-dependent dilation as assessed by FMD of the brachial artery was evident after only 2 weeks, with further increases after 4 and 8 weeks.36 In hypercholesterolemic patients with perfusion abnormalities, 12 weeks of treatment with fluvastatin (40 to 80 mg/d) significantly increased myocardial perfusion in ischemic segments by 30% (P<0.001). In normal segments, perfusion increased by only 5% (P<0.005).37 Another study in patients with moderately elevated cholesterol levels showed that the vasodilator response to acetylcholine as determined by forearm blood flow was significantly (P<0.005) increased after 1 month of treatment with simvastatin 20 mg/d, and this improvement was further enhanced after 3 months.38 In the Reduction of Cholesterol in Ischemia and Function of the Endothelium (RECIPE) trial, 6 weeks of pravastatin therapy (40 mg/d) rapidly increased FMD compared with placebo in patients with acute coronary syndromes. Changes in FMD were not correlated with decreases in total and LDL cholesterol, suggesting that the improvement in endothelial function was not related to the lipid-lowering effects of the statin.39 Augmented endothelium-dependent dilation has been noted in the forearm of healthy, normocholesterolemic men after only 1 day of high-dose atorvastatin (80 mg), even before appreciable reduction in plasma LDL cholesterol or CRP could be detected. This rapid increase in dilation is consistent with a cholesterol-independent effect of statins.40

However, a study in patients with CAD and mildly elevated cholesterol levels found no difference at 6 months in coronary endothelial vasomotor function between placebo and simvastatin (40 mg/d) groups, although simvastatin therapy markedly improved the lipid profile.41 Endothelial function was assessed in both epicardial arteries (using acetylcholine and angiography) and the microvasculature (coronary blood flow response to substance P). The authors proposed several explanations for the discrepancy between their findings and those of other studies. For example, subjects in this study had lower baseline total and LDL cholesterol levels, less severe baseline atherosclerosis, and relatively mild baseline endothelial dysfunction compared with subjects in studies in which improved endothelial function was observed.42

Identifying the Mechanism
Improvement in endothelium-dependent vasodilation has been achieved with cholestyramine and LDL-apheresis, implicating LDL cholesterol reduction as an important mechanism. However, the clinical benefits observed with statin therapy in large controlled trials appear to exceed the benefits that would be expected from the observed reductions in cholesterol, suggesting that nonlipid effects (as part of their multiple effects referred to as pleiotropic effects), including improvement in endothelial dysfunction, may be involved.44 A number of mechanisms have been proposed for these effects. One important pathway appears to be the effects of statins on NO production via increased availability of eNOS.

In one study in hypercholesterolemic patients, the significant improvement in endothelium-dependent vasodilation noted after statin therapy was blocked by administration of L-NMMA, an inhibitor of eNOS. This finding suggested that increased bioavailability of NO was the mechanism for the improved endothelial function.22 In another study, short-term therapy with pravastatin improved forearm vasodilation in normocholesterolemic subjects with CAD.45 This improve-
ment was likewise blocked by the coadministration of l-NMMA. Of note, the improvement in endothelial vasodilatory function was independent of the cholesterol-lowering effects of statin therapy. As observed previously, the rapid action of statins that precedes a reduction in serum cholesterol is also suggestive of a lipid-independent effect.40

Elevated levels of native LDL decrease the bioavailability of endothelium-derived NO and downregulate endothelial eNOS. A recent study using endothelial cells from human umbilical veins showed that simvastatin prevented the downregulation of eNOS by native LDL.46 Elevated LDL reduces NO production in part by increasing the interaction between caveolin-1 and eNOS.47 An in vitro study has shown that atorvastatin reduces caveolin-1 expression in endothelial cells, inhibiting the interaction between caveolin-1 and eNOS and resulting in increased NO production. The statin-induced reduction in caveolin-1 expression was independent of the extracellular LDL-cholesterol level.47 The statin also promoted the agonist-induced association of eNOS and the chaperone heat shock protein 90 (Hsp90), resulting in the potentiation of eNOS activation.47 Another proposed mechanism for the increase in eNOS expression with statin therapy is increased stability of eNOS messenger RNA, which would permit preservation of eNOS expression in the presence of oxidized LDL.48

Activation of the Rho/Rho kinase pathway reduces the stability of eNOS mRNA, whereas inhibition of this pathway augments the stability of eNOS mRNA. Statins inhibit the activity of Rho/Rho signaling by blocking the generation of geranylgeranyl pyrophosphate, an effect that is cholesterol-independent.49

Statin therapy also reduces circulating levels of the adhesion molecules P-selectin and intercellular adhesion molecule-1 (ICAM-1) in hypercholesterolemic subjects.50 The reduction in adhesion molecules was associated with an increase in levels of NO.50 These findings suggest that statins may reduce platelet and leukocyte adhesion as well as improve endothelial cell function.50

The serine/threonine protein kinase Akt (protein kinase B) mediates the activation of eNOS, resulting in increased production of NO. Statins have been found to activate Akt in endothelial cells, thereby enhancing the phosphorylation of the endogenous Akt substrate eNOS and producing an increase in NO.51

Oxidative stress has been shown to contribute to endothelial dysfunction and the development of atherosclerosis. The use of antioxidant supplementation to improve endothelial function has support from some studies. For example, in one study, an LDL-lowering plus antioxidant regimen (lovastatin plus probucol) improved endothelium-dependent vasomotor response.52 In another study, probucol was found to be useful in preventing coronary restenosis, whereas antioxidant vitamins failed to exert any benefit.53 Antioxidant vitamins have not been found beneficial in several clinical trials including the recent Heart Protection Study.54 However, these findings should not be taken as the final word on the use of antioxidants in CHD patients.10 Antioxidants are a diverse group of compounds, and in view of the central role played by oxidized LDL in endothelial dysfunction and atherogenesis, the antioxidant effects of statin therapy and probucol-like compounds are still of great therapeutic interest. Statins prevent the downregulation of eNOS by oxidized LDL.48 Finally, statins also appear to reduce the atherogenic potential of lipoproteins. Atorvastatin hydroxy metabolites inhibit oxidation of LDL, VLDL, and HDL.55 Atorvastatin has also been shown to suppress the cellular uptake of oxidized LDL by monocytes, a key step in the development of atherosclerosis.56

Conclusions

Endothelium and its product nitric oxide are key regulators of vascular health. Reduced bioavailability of NO is involved in the initiation, progression and complications of atherosclerosis. Not surprisingly, deficiency of nitric oxide in coronary or peripheral arteries is predictive of future cardiovascular events. Basic mechanisms involved in endothelial dysfunction have been clarified and suggest a plethora of new therapeutic targets. In particular, studies investigating the nonlipid effects of statins on vascular function constitute a promising avenue of research.

References

19. Reddy KG, Nair RN, Sheehan HM, et al. Evidence that selective endothelial dysfunction may occur in the absence of angiographic or
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