Carbon Monoxide and Nitric Oxide Mediate Cytoskeletal Reorganization in Microvascular Cells via Vasodilator-Stimulated Phosphoprotein Phosphorylation: Evidence for Blunted Responsiveness in Diabetes

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OBJECTIVE—We examined the effect of the vasoactive agents carbon monoxide (CO) and nitric oxide (NO) on the phosphorylation and intracellular redistribution of vasodilator-stimulated phosphoprotein (VASP), a critical actin motor protein required for cell migration that also controls vasodilation and platelet aggregation.

RESEARCH DESIGN AND METHODS—We examined the effect of donor-released CO and NO in endothelial progenitor cells (EPCs) and platelets from nondiabetic and diabetic subjects and in human microvascular endothelial cells (HMECs) cultured under low (5.5 mmol/l) or high (25 mmol/l) glucose conditions. VASP phosphorylation was evaluated using phosphorylation site-specific antibodies.

RESULTS—In control platelets, CO selectively promotes phosphorylation at VASP Ser-157, whereas NO promotes phosphorylation primarily at Ser-157 and also at Ser-239, with maximal responses at 1 min with both agents on Ser-157 and at 15 min on Ser-239 with NO treatment. In diabetic platelets, neither agent resulted in VASP phosphorylation. In nondiabetic EPCs, NO and CO increased phosphorylation at Ser-239 and Ser-157, respectively, but this response was markedly reduced in diabetic EPCs. In endothelial cells cultured under low glucose conditions, both CO and NO induced phosphorylation at Ser-157 and Ser-239; however, this response was completely lost when cells were cultured under high glucose conditions. In control EPCs and in HMECs exposed to low glucose, VASP was redistributed to filopodia-like structures following CO or NO exposure; however, redistribution was dramatically attenuated under high glucose conditions.

CONCLUSIONS—Vasoactive gases CO and NO promote cytoskeletal changes through site- and cell type–specific VASP phosphorylation, and in diabetes, blunted responses to these agents may lead to reduced vascular repair and tissue perfusion. 

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The gaseous signal molecules nitric oxide (NO) and carbon monoxide (CO) exert multiple modulatory actions in regulating vascular function. While NO effects have been recognized for over a decade, similar vasoregulatory action of CO was established only recently. CO is generated by heme oxygenase (HO)-1 under a wide variety of conditions (e.g., cell exposure to such stressors as hypoxia, growth factors, and cytokine stimulation) that activate the enzyme (1,2). Unlike its highly reactive cognate NO, which participates in multiple redox reactions, CO is a relatively stable gas that exhibits extraordinary affinity for heme centers (3–5). Like NO, the signaling effects of CO rely in part on its ability to form a complex with the heme moiety of soluble guanylate cyclase (sGC), stimulating the synthesis of the diffusible second messenger guanosine 3’5’-cyclic monophosphate (cGMP) (6). The sGC/cGMP pathway plays a critical role in mediating the effects of CO on vascular relaxation and inhibition of platelet aggregation and coagulation (7,8). A recently recognized property of NO is its cell type–specific facilitation or inhibition of cell migration (9), a complex process involving molecular-mechanical events that depend on extracellular signaling, actin-based motility, and cell adhesion. Endothelial progenitor cells (EPCs) differentiate into endothelial cells whose function in vascular repair depends on chemokine- and growth factor–directed cell migration. The role of EPCs in endothelial repair is supported by their ability to inhibit development of atherosclerosis (10,11) and intimal hyperplasia (12), while still promoting beneficial angiogenesis. We previously demonstrated the central role of the actin cytoskeleton in EPC migration (13), and our findings suggest that NO has a critical function within EPCs, where it regulates the distribution of vasodilator-stimulated phosphoprotein (VASP). The latter plays a pivotal role in promoting actin filament elongation at the leading edge by forming an active molecular motor complex that propels motility (14). VASP contains three distinct phosphorylation sites (Ser-157, Ser-239, and Thr-278), the first of which is preferentially phosphorylated by cAMP-dependent protein kinase (PKA) and the second by cGMP-dependent protein kinase (PKG). Although the exact roles of phosphorylated residues in VASP have not completely been elucidated, one idea is that a high 3’,5’-cyclic AMP (cAMP)-to-cGMP ratio promotes VASP-activated actin filament elongation, whereas a low cAMP-to-cGMP ratio favors filament capping and loss of motility (15). The following factors are...
known to influence VASP phosphorylation: intracellular localization in focal adhesions, filopodia, and lamellipodia; accessibility of phosphorylation sites in VASP that is complexed with other proteins; availability of specific protein kinases and/or phosphoprotein phosphatases; and the respective activators and inhibitors of these kinases and phosphatases (16). We previously reported that the reduced bioavailability of NO in diabetic individuals prevents VASP redistribution, resulting in the inability of EPCs to form proper cytoskeletal extensions (13). We also showed that the EPC chemoattractant stromal cell–derived factor-1 (SDF-1) transcriptionally activates HO-1 via the atypical protein kinase C (PKC)-ζ isoform generating CO, which in turn can phosphorylate VASP in endothelial cells (17).

Because PKG and PKA catalyze VASP phosphorylation, and because the latter is thought to control VASP’s subcellular distribution and function (13), we directly compared the effects of NO and CO on VASP phosphorylation and redistribution in cells typically known to be dysfunctional in diabetes, namely platelets, EPCs, and microvascular endothelial cells. We demonstrate that both CO and NO regulate VASP phosphorylation and that pretreatment with either agent stimulates migration toward SDF-1. We also show that normal platelets display a modest response to exogenous CO stimulation but a greater response to NO treatment. In contrast, diabetic platelets are not responsive to either CO or NO treatment (data not shown). Culturing microvascular endothelial cells at high glucose concentrations also results in reduced VASP phosphorylation. These novel findings suggest that CO regulates VASP phosphorylation and vascular cell migration under conditions of reduced NO bioavailability, as observed in diabetes.

**RESEARCH DESIGN AND METHODS**

**Cell culture.** Human lung microvascular endothelial cells (HMVECs) (LONZA, Walkersville, MD) were cultured to 80% confluency in EGM-2 MV medium (Lonza) in a humidified 5% CO2/95% air atmosphere. Cells were incubated overnight in EBM-2 supplemented with 2% fetal bovine serum. Fresh medium (EBM-2 + 2% fetal bovine serum) was added on the following day, and CO- or NO-containing medium was added to the designated samples for 1 or 15 min.

**Isolation of platelets.** Peripheral blood from healthy and diabetic volunteers was then collected in cell preparation tubes (CPTs) with heparin (BD Biosciences) and centrifuged to obtain the mononuclear cell fraction, from which CD34+ cells were isolated using a CD34+ isolation kit (Stem Cell Technologies, Vancouver, CA) according to the manufacturer’s protocol, as previously described (13).

**Immunohistochemistry.** EPCs were cultured on fibronectin-coated dishes with Endocult stem cell liquid media until colonies formed. These cells were then treated for 15 min or 4 h with 100 μmol/l diethylenetriamine (DETA)-NO (Sigma-Aldrich, St. Louis, MO) in water or 10 μmol/l NO-containing medium was added to the designated samples for 1 or 15 min (Stem Cell Technologies, Vancouver, CA). Briefly, 2 × 10^7 cells were incubated with a DETA-NO selection cocktail for 15 min, and 50 μl magnetic nanoparticles was then added and incubated for another 10 min. The cell suspension volume was increased to 2.5 ml, and the particle-bound CD34+ cells were concentrated on the tube’s inner surface with the aid of a magnet. The supernatant was decanted, and the remaining CD34+ cells were resuspended in RPMI culture media (Cellgro, Herndon, VA).

**Fluorescence-activated cell sorting.** Peripheral blood was collected into CPTs (BD Biosciences) and centrifuged to obtain the mononuclear cell fraction, from which CD34+ cells were isolated using a CD34+ isolation kit (Stem Cell Technologies, Vancouver, CA). Briefly, 2 × 10^7 cells were incubated with a DETA-NO selection cocktail for 15 min, and 50 μl magnetic nanoparticles was then added and incubated for another 10 min. The cell suspension volume was increased to 2.5 ml, and the particle-bound CD34+ cells were concentrated on the tube’s inner surface with the aid of a magnet. The supernatant was decanted, and the remaining CD34+ cells were resuspended in RPMI culture media (Cellgro, Herndon, VA).

**Preparation of DETA-NO and Ru(II)(CO)3Cl2 donors.** The half-lives for NO and CO release from DETA-NO and Ru(II)(CO)3Cl2 are greatly different, resulting in the almost immediate release of CO but very slow release of NO. Therefore, to assure that cells were exposed to comparable concentrations of NO and CO, we first calculated the amount of CO released at 1 and 15 min using the integrated rate law: [CO donor] / [CO donor]_irr = exp (-k_CO), where [CO donor]_irr is the CO donor concentrations at time t, and k_CO is the first-order rate constant for CO release, and t is the incubation period. We then used the corresponding rate law, i.e., [NO donor] / [NO donor]_irr = exp (-k_NO), to determine the time needed to obtain an extent of NO release comparable with that observed with CO donor in 1 and 15 min. The CO and NO donor rate constants k_CO and k_NO were derived from the rate equation: k_CO = [CO donor] / [CO donor]_irr = exp (-k_CO) for 1 min, followed by two additional 10-min washes with TBS. Protein bands were reprobed for VASP using a commercially available BCA kit (Pierce, Rockford, IL), and 50 μg was applied to each well above the polyacrylamide gel (7.5%) (Criterion; Biorad Laboratories, Richmond, CA) at 120 V for 20 min, followed by 140 V for 65 min. Samples were run in duplicate for later detection of the two phosphorylated VASP (pVASP) isoforms. The separated proteins were transferred to a nitrocellulose membrane (Biorad Laboratories) in a semidry transblot apparatus (Biorad Laboratories). The membranes were blocked with 5% milk and 1% Tween-20 (Sigma-Aldrich) for 1 h at room temperature. For the detection of pVASP isoforms, the membranes were incubated at 4°C overnight with either a mouse monoclonal anti-VASP Ser-239 antibody (Upstate Cell Signaling Solutions, Lake Placid, NY) or a rabbit polyclonal anti-VASP Ser-157 antibody (Santa Cruz Biotechnology, Santa Cruz, CA), both at a 1:200 dilution. Blots were then washed with 5% milk for 5 min and incubated with an HRP-conjugated anti-rabbit antibody (Sigma-Aldrich) or an anti-rabbit (Santa Cruz Biotechnology) for 1 h at room temperature. After incubation with the secondary antibody, the membranes were washed twice with TBS containing 0.05% Tween-20 for 5 min, followed by two additional 10-min washes with TBS. Protein bands were visualized with enhanced chemiluminescence using a commercial Western blot detection kit (Amersham Biosciences, Amersham, U.K.) and pVASP levels were analyzed using ImageQuant software Version 2.1 (1) for Standard. Protein bands were analyzed by comparing with molecular weight markers (Biorad Laboratories) served to verify the molecular mass of pVASP (50 kDa).

After pVASP detection, the membranes were probed for VASP using a 1:200 dilution of a goat anti-VASP antibody (Santa Cruz Biotechnology) and...
for β-actin using a mouse anti-β-actin primary antibody at a dilution of 1:7,000 (Sigma-Aldrich), followed by HRP-conjugated anti-goat (Santa Cruz Biotechnolog) and anti-mouse IgG antibody (Sigma-Aldrich), both at 1:2,000 dilution, and were visualized by enhanced chemiluminescence. Standard molecular weight markers served to verify the 42-kDa molecular mass of β-actin.

**In-cell Western analysis.** CD34\(^{+}\) cells were cultured in defined serum-free medium (StemSpan SFEM, StemCell Technologies, Vancouver, Canada) to obtain an optimal number of cells. For CD34\(^{+}\) cell proliferation and expansion, without differentiation, we used 1 ml StemSpan SFEM supplemented with a cytokine cocktail (10 ng/ml IL-3, 100 ng/ml stem cell factor, 20 ng/ml interleukin-3, and 20 ng/ml interleukin-6, StemCell Technologies) and 50 ng/ml thrombopoietin (R&D Systems, Minneapolis, MN). After determining cell number with a hemocytometer (Hauser Scientific, Horsham, PA), 10,000 cells were transferred to each well in a 96-well plate (BD Falcon, San Jose, CA). After incubation with/without CO or NO donor, cells were fixed with 4% paraformaldehyde for 20 min at room temperature and then harvested by centrifugation. After removing the fixation solution, cells were permeabilized with 0.1% Triton X-100 in PBS and blocked with LI-COR Odyssey blocking buffer (LI-COR Biosciences, Lincoln, NE) for 1.5 h at room temperature with moderate shaking. Blocking buffer was removed by aspiration, and the cells were incubated with 50-μl volume of diluted primary antibody at 4°C overnight in a cold room. Cells were then washed four times for 5 min in PBS containing 0.1% Tween-20 (Fisher Scientific, Pittsburgh, PA) and incubated with diluted secondary antibody. After 1-h incubation, cells were washed four times (5 min each) with PBS containing 0.1% Tween-20, followed by final washing in PBS to remove excessive detergent. The 96-well plate was then scanned in the appropriate channels using an Odyssey infrared imaging system (LI-COR Biosciences). Relative quantification was normalized and readjusted in cell number from well to well using DNA staining.

**Statistical analysis.** All reported data are expressed as the mean ± SEM and were obtained from at least three independent experiments. Student’s t test and ANOVA with Student-Newman-Keuls posttest were used to compare the data.

**RESULTS**

The choice of the technique used (e.g., standard Western blotting, In-Cell Western analysis, and FACS) for these studies was largely governed by the number of cells available for analysis. Because CD34\(^{+}\) cells represented <1% of circulating mononuclear cells, In-Cell Western (low sensitivity) and FACS (higher sensitivity) analyses were the methods of choice; however, for more abundant cells such as endothelial cells, standard Western blotting was sufficient. All the phosphorylation studies on EPCs were performed after 15 min of treatment. We then tested whether a shorter treatment (1 min) could still elicit a significant migratory response.

We therefore pretreated human CD34\(^{+}\) EPCs for 1 and 15 min with medium containing NO or freshly generated CO, followed by SDF-1 stimulation (13), and then assessed cell migration using Boyden chambers. As demonstrated in Fig. 1, both CO and NO treatment resulted in increased EPC migration to SDF-1, although a more robust response was observed with NO at 1 min.

Because VASP is a key component of the actoclampin molecular motors responsible for actin-based cell motility (14,20), and because VASP was originally identified as a major substrate for cGMP-stimulated protein kinase in platelets and endothelial cells (21), we investigated whether NO and CO elicited similar effects on VASP phosphorylation and its mobilization to the leading edge of motile cells. VASP function appears to be chiefly regulated by phosphorylation at Ser-157 and Ser-239, and the availability of phosphorylation site-specific antibodies provides an unambiguous way to define both the sites and extents of signal molecule–activated VASP phosphorylation. We therefore incubated CD34\(^{+}\) cells for 15 min in the presence of NO or CO already liberated from their respective donors. As shown in Fig. 2, NO treatment routinely increased the extent of VASP phosphorylation at Ser-239, whereas CO treatment consistently increased the extent of Ser-157 phosphorylation in these cells.

To confirm that the differential effects of CO and NO are not limited to CD34\(^{+}\) cells, we also examined platelets, which are a rich source of VASP and are the cells wherein VASP phosphorylation is best characterized (22). In platelets exposed to either NO or CO for 1- and 15-min periods, we found that CO treatment caused a modest yet significant VASP phosphorylation at Ser-157, whereas NO treatment significantly increased VASP phosphorylation at Ser-157 with maximum response at 1 min and at Ser-239 with maximum effect at 15 min (Fig. 3). These findings suggest that CO and NO elicit markedly different VASP phosphorylation responses in platelets and CD34\(^{+}\) cells.
Because VASP resides on the cytoplasmic face of the leading edge in motile cells (23), and because VASP phosphorylation is believed to control its interactions with VASP-docking proteins like vinculin, zyxin, and migfilin, we next examined the effects of donor-generated NO and CO on the redistribution of VASP in microvascular endothelial cells growing while firmly adhered to fibronectin-coated culture dishes. Under basal conditions (i.e., no NO or CO), VASP was mainly localized along the sides of actin filaments found throughout the peripheral cytoplasm (Fig. 4A and B). However, following a 15-min exposure to CO, VASP redistributed to the leading edge of advancing microvascular endothelial cells (Fig. 4C and D). VASP redistribution was also observed when these cells were incubated with NO (Fig. 4E and F).

In as much as diabetes-associated vascular dysfunction is frequently attributed, at least in part, to reduced levels of bioavailable NO, we sought to determine whether diabetes impacted the NO and CO effects on VASP phosphorylation within cell types typically affected by diabetes. We treated platelet samples from diabetic individuals with NO or CO under the same conditions used in Fig. 1 and observed no change in VASP phosphorylation, at either Ser-157 or Ser-239 (data not shown). In Fig. 5, VASP phosphorylation is shown for two diabetic patients and one nondiabetic control subject. Patient 1 had type 2 diabetes of 5 years duration and excellent glycemic control (A1C 6.4%), while patient 2 had type 1 diabetes of 48 years duration and poor glycemic control (A1C 11.3%). A different pattern was observed in the diabetic CD34+ cells, where NO treatment stimulated phosphorylation at VASP's Ser-239 to a significantly less extent, when compared with the nondiabetic control cells. However, we observed considerable patient-to-patient variation in the degree of the response but all with the same pattern.

Notably, compared with control cells grown in low-glucose medium, exposure of microvascular cells to conditions mimicking aspects of diabetes (e.g., culturing these cells for 1 week in high-glucose medium) resulted in a failure of NO and CO to elicit changes in either VASP redistribution to the leading edge of the cells, as shown by immunofluorescence (compare Figs. 4 and 6), or phosphorylation at Ser-157 or Ser-239, as detected by standard Western blot analysis (Fig. 7).

**Discussion**

Understanding the regulation of endothelial cell migration in response to chemokines has proven to be a daunting task that requires the investigation of a manifold of factors known to affect chemotaxis (e.g., chemokine sensing by receptor-mediated signaling, cellular locomotion by actin-based molecular motors, and even the cell's energy status). Previous studies from our laboratory on CD34+ EPCs showed that diabetic EPCs have reduced intracellular NO concentration as well as a concomitantly reduced migratory capability. We found that, when pretreated with an NO donor, cell migration can be restored (13), and we further demonstrated that this exogenous NO exposure resulted in enhanced diabetic cell migration, with attendant changes in phosphorylation of the actin cytoskeletal protein VASP (13).

Given the robust nature of the vasodilator gases CO and NO in modulating vascular function, we focused on the following questions: Can CO suffice in place of NO in promoting vascular cell migration? If so, do NO and CO exert their signaling effects on the same downstream phosphorylation target(s)? In this study, we demonstrate that CO can regulate VASP phosphorylation and in turn alter cell migration. We also directly compare the CO effects to those of NO in EPCs, platelets, and microvascular endothelial cells. Our studies demonstrate that both vasoactive gases promote cytoskeletal changes through site- and cell type-specific VASP phosphorylation; however, these responses to NO and CO are blunted in diabetes and may be responsible for reduced vascular repair and tissue perfusion (24,25).

The actoclampins are the force-generating motors responsible for actin-based cell crawling and vesicle motility (14). Each of these membrane surface–bound molecular motors consists of an actin filament plus-end tracking protein called a clampin (e.g., VASP, N-WASP [neural Wiskott–Aldrich syndrome protein], formins, etc.) and its actin filament partner. The energy for force production
appears to be derived from nucleotide hydrolysis at the filament’s penultimate actin-ATP subunits, thereby promoting clamping release, translocation, and rebinding to terminal actin-ATP subunits. Before the active motor complex is assembled, clampins must be recruited to the membrane surface, where they dock at motility sites at the tips of filopodia and lamellipodia as well as in the focal adherens complex. Rottner et al. (23), for example, showed that VASP not only colocalizes to adhesion sites with the adaptor proteins vinculin and zyxin, but is also recruited to the tips of lamellipodia in amounts that are directly proportional to the rate of lamellipodial protrusion. Tokuo and Ikebe (26) further showed that myosin X specifically transports VASP and other members of the Ena/VASP clamping protein family to the leading edge, where VASP then binds to a membrane docking protein such as vinculin, zyxin, or migfilin (27,28). Only then can the actoclampin motor assemble and generate the forces needed for cell migration. When viewed from this perspective, VASP phosphorylation may affect the recruitment by myosin X, VASP docking with membrane components, and/or VASP-mediated formation of an active motor that can propel cell crawling. Moreover, as cells stop moving, VASP is known to redistribute to other intracellular sites. Benz et al. (29), for example, showed that VASP interacts with αII-spectrin in endothelial cells and that PKA-mediated VASP phosphorylation at Ser-157 inhibits this binding interaction. They also showed that VASP is dephosphorylated upon formation of cell-cell contacts and that, in confluent cells, αII-spectrin colocalizes with nonphosphorylated VASP at cell-cell junctions (29). The exact details of how VASP phosphorylation at Ser-157 or Ser-239 controls one or more of these steps remain to be worked out. Even so, the observation that different cell types contain various clampins and their respective membrane-docking proteins is likely to explain why VASP phosphorylation can either stimulate or suppress cell crawling and actin-based cell shape changes in a manner depending on cell type and/or culture conditions.

VASP is also the most abundant platelet and endothelial protein phosphorylated by PKG in NO signaling pathways, and, as noted above, this cytoskeletal adaptor protein is also a PKA substrate. Three serine/threonine phosphorylation sites within VASP play roles resulting in the inhibition of platelet aggregation and focal adhesion assembly (30,31). In this study, we demonstrated that CO can also regulate VASP phosphorylation, and we directly compare CO and NO effects in EPCs, platelets, and microvascular endothelial cells. We found that CO pretreatment can similarly stimulate EPC migration (Fig. 1) and phosphor-

![Image](https://example.com/image.png)

**FIG. 4.** CO- and NO-mediated VASP redistribution in HMECs. HMECs, cultured on fibronectin-coated coverslips, were treated with CO and NO donors as described in the legend to Fig. 1. A and B: Low- and high-power images of untreated cells showing uniform VASP localization on actin filaments throughout the cytoplasm. C and D: Low- and high-power images showing CO-induced VASP redistribution to filopodia at the leading edge of microvascular endothelial cells. E and F: Low- and high-power images showing NO-induced redistribution of VASP to filopodia. Representative results from three independent experiments are shown. Green, VASP; blue, DAPI (nuclei). Scale bars = 25 μm. (Please see http://dx.doi.org/10.2337/db08-0381 for a high-quality digital representation of this figure.)

| % VASP Ser-239 phosphorylation (relative to Control at time 0) |
|-----------------|-----------------|-----------------|
| 0 hour | 0.25 hour | 4 hour |
| Control | DA | DB |

**FIG. 5.** NO exposure results in reduced VASP Ser-239 phosphorylation in diabetic CD34+/H11001 cells compared with control cells. The extent of phosphorylation was determined by FACS analysis on CD34+/H11001 cells from two diabetic subjects (DA [diabetic donor A] and DB [diabetic donor B]) and cells from a nondiabetic subject (Control [normal donor]). Values represent means ± SD. *P < 0.05.

While we did not perform direct measurements of CO levels in our experiments, we did use CO concentrations that are achievable in vivo in physiological conditions and previously used by investigators (32,33). Endogenous CO has been reported to be generated in many cell types, and the amount of CO released via the heme oxygenase reaction can reach up to 12 ml/day (~16 μmol·1·h⁻¹) (32). Previous studies have reported that tissues can produce 0.1–100 μmol/l CO in vivo from the HO reaction (33); it is therefore quite reasonable to believe that sufficient levels of HO-1 are present in cells to provide sufficient levels of CO, explaining our observed changes in VASP phosphorylation.
Moreover, in response to incubation of microvascular endothelial cells with NO or CO donors, VASP was readily redistributed to the peripheral membrane and filopodia (Fig. 4). Thus, with regard to VASP localization, our data suggest that CO and NO may both have critical roles in vascular cell dynamics, and CO may have important contributions at low NO bioavailability, a condition already known to occur in diabetes. Our studies clearly indicate that both vasoactive gases promote cytoskeletal changes through site- and cell type-specific VASP phosphorylation and that responses to NO and CO are blunted in diabetes. While the migratory deficiency seen in diabetic EPCs could be overcome by exogenous NO (13), exogenous NO did not result in phosphorylation and redistribution of VASP in mature endothelial cells cultured in high glucose conditions. This defect can be viewed as “diabetes-induced NO resistance.”

What also becomes apparent from the present study is that exposure of cells to NO or CO can greatly alter VASP recruitment to the leading edge with consequential effects on cell motility. Perhaps even more significant, in the context of diabetes, is that culturing endothelial cells at high glucose conditions results in motility defects that can be traced, at least in part, to altered VASP phosphorylation. We therefore postulate that these defects contribute to reduced vascular repair and tissue perfusion.

Both heme oxygenases generate CO, but they do so with very different kinetics (34,35); HO-1 is induced by oxidants such as hydrogen peroxide, UV radiation, and proinflam-
matory cytokines and by growth factors, hemodynamic or shear stress, heat shock, and even by NO (3). Endothelial cells derived from either the micro- or macrovasculature (36) responded equally to CO and NO, reflecting the key roles of both HO-1 and endothelial NO synthase throughout in the entire vasculature. In contrast, platelets do not respond to CO, perhaps reflective of the low levels of heme oxygenase in adult platelets (37). By altering HO-1 and NOS gene expression, hypoxia potentially modulates the availability of these gaseous second messengers. Thus, fluxes in the CO and NO generation during hypoxic stress are likely to have dramatic consequences on the regulation of vascular functions such as dilation, expression of vasomodulators, inhibition of plateau aggregation, and smooth muscle cell proliferation (38). Although the findings in this report support a role for CO sufficing for NO in promoting vascular cell migration, the signaling action of CO results in the phosphorylation of different VASP sites than NO. Furthermore, we demonstrate that, while both vasoactive gases promote cytoskeletal changes through site- and cell type–specific VASP phosphorylation, these responses are blunted in diabetes and may be responsible for the altered vascular repair and tissue perfusion associated with diabetic vascular complications.

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