Nitric oxide alterations following acute ductal constriction in the fetal lamb: a role for superoxide

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Hsu J, Oishi P, Wiseman DA, Hou Y, Chikovani O, Datar S, Sajti E, Johengen MJ, Harmon C, Black SM, Fineman JR. Nitric oxide alterations following acute ductal constriction in the fetal lamb: a role for superoxide. Am J Physiol Lung Cell Mol Physiol 298: L880–L887, 2010. First published April 2, 2010; doi:10.1152/ajplung.00384.2009.—Acute partial compression of the fetal ductus arteriosus (DA) results in an initial abrupt increase in pulmonary blood flow (PBF), which is followed by a significant reduction in PBF to baseline values over the ensuing 2–4 h. We have previously demonstrated that this potent vasoconstricting response is due, in part, to an endothelin-1 (ET-1)-mediated decrease in nitric oxide synthase (NOS) activity. In addition, in vitro data demonstrate that ET-1 increases superoxide levels in pulmonary arterial smooth muscle cells and that oxidative stress alters NOS activity. Therefore, the objectives of this study were to determine the potential role of superoxide in the alterations of hemodynamics and NOS activity following acute ductal constriction in the late-gestation fetal lamb. Eighteen anesthetized near-term fetal lambs were instrumented, and a lung biopsy was performed. After a 48-h recovery, acute constriction of the DA was performed by inflating a vascular occluder. Polyethylene glycol-superoxide dismutase (PEG-SOD; 1,000–1,500 units/kg, n = 7) or PEG-alone (vehicle control group, n = 5) was injected into the pulmonary artery before ductal constriction. Six animals had a sham operation. In PEG-alone-treated lambs, acute ductal constriction rapidly decreased pulmonary vascular resistance (PVR) by 88%. However, by 4 h, PVR returned to preconstriction baseline. This vasoconstriction was associated with an increase in lung superoxide levels (82%), a decrease in total NOS activity (50%), and an increase in P-eNOS-Thr495 (52%) (P < 0.05). PEG-SOD prevented the increase of superoxide after ductal constriction, attenuated the vasoconstriction, preserved NOS activity, and increased P-eNOS Ser1177 (307%, P < 0.05). Sham procedure induced no changes. These data suggest that an acute decrease in NOS activity that is mediated, in part, by increased superoxide levels, and alterations in the phosphorylation status of the endothelial NOS isoform, underlie the pulmonary vascular response to acute ductal constriction.

ductus arteriosus; pulmonary circulation

INCREASES IN Fetal PULMONARY arterial pressure induced by mechanical constriction of the ductus arteriosus induce an acute increase in pulmonary blood flow that is followed by active vasoconstriction (1). This so-called “myogenic response,” which returns pulmonary blood flow to preconstriction values within 2–4 h, may represent an adaptive response of the fetal pulmonary vasculature to maintain the normal low flow state (24). However, chronic ductal constriction results in pulmonary vascular remodeling and many of the pathophysiological features of persistent pulmonary hypertension of the newborn (2, 16, 31). In fact, fetal ductal constriction secondary to maternal use of COX inhibitors, such as indomethacin, has been associated with persistent pulmonary hypertension of the newborn (26). Therefore, understanding the mechanisms that mediate changes in pulmonary blood flow following ductal constriction has important clinical implications.

Increasing evidence demonstrates that factors produced by the pulmonary vascular endothelium regulate normal fetal pulmonary vascular tone, as well as the dynamic changes in pulmonary vascular tone following both acute and chronic constriction of the ductus arteriosus. For example, both acute and chronic ductal constriction are associated with decreased nitric oxide (NO) activity and increased endothelin-1 (ET-1)-mediated vasoconstriction (6, 13, 17, 22, 27). In addition, further studies suggest that this decrease in NO activity mediates, in part, the hemodynamic response following acute ductal constriction and are independent of changes in gene expression and dependent on ET-1 activity (12, 17, 23). Moreover, increasing data demonstrate that changes in the redox environment may alter NOS activity by several mechanisms including changes in the nitration and phosphorylation status of the enzyme (5). Last, ET-1 has been demonstrated to increase superoxide levels in fetal pulmonary arterial endothelial cells (19). However, the potential role of these superoxide-NO interactions in mediating the acute changes in fetal pulmonary blood flow following ductal constriction has not been investigated.

Therefore, the objective of this study was to determine the role of superoxide, NO, and their interactions in regulating the dynamic changes in fetal pulmonary blood flow following acute mechanical constriction of the ductus arteriosus. We hypothesized that acute mechanical constriction of the ductus arteriosus would result in increased superoxide production, and superoxide-dependent decreases in NOS activity, with a net result of active pulmonary vasoconstriction. To investigate this hypothesis, we determined superoxide levels, NOS activity, and posttranslational modifications of NOS, before and 4 h after mechanical ductal constriction in late-gestation fetal lambs. To determine potential superoxide-NO interactions following ductal constriction, the same factors were studied in an additional group of fetal lambs that were pretreated with polyethylene glycol-superoxide dismutase (PEG-SOD), a superoxide scavenger. Last, to isolate changes related to the experimental protocol, we studied the same factors in an additional group of fetal lambs that underwent a sham procedure without ductal constriction.
MATERIALS AND METHODS

Surgical preparation. Eighteen mixed-breed Western pregnant ewes (132–140 days’ gestation, term = 145 days) were operated on under sterile conditions with the use of local (2% lidocaine hydrochloride) and intravenous anesthesia (0.002 mg·kg⁻¹·min⁻¹ diazepam and 0.3 mg·kg⁻¹·min⁻¹ ketamine hydrochloride). Fetal anesthesia consisted of local anesthesia with 2% lidocaine hydrochloride and 15 mg/kg im ketamine hydrochloride. Through a uterine incision, the fetal forelimb was exposed. Polyvinyl catheters were inserted into the fetal pedal artery and vein and were advanced to the aorta and the inferior vena cava, respectively. A left lateral thoracotomy was performed in the fourth intercostal space. The pericardium was incised along the main pulmonary trunk. Teflon cannulas attached to polyvinyl catheters were inserted into the proximal main pulmonary trunk, left pulmonary artery, and the left atrium. An ultrasonic flow transducer (Transonic Systems, Ithaca, NY) was placed around the left pulmonary artery. The ductus arteriosus was dissected free and infiltrated with 10% formalin to prevent ductal constriction during manipulation. A vascular occluder was then placed around the ductus arteriosus, but left uninflated. A side-biting vascular clamp was utilized to isolate peripheral lung tissue from the right upper lobe, and the incision was cautereized. Approximately 300 mg of peripheral lung were obtained for the biopsy to determine NOS activity, endothelial nitric oxide synthase (eNOS) protein levels, and superoxide levels. The thoracotomy incision was then closed in layers. Warm saline was instilled to replace the lost amniotic fluid, and the uterine incision was closed. A polyvinyl catheter was placed in the amniotic cavity. The catheters were filled with PBS containing 1,000 U/ml heparin sodium, plugged, and brought to the skin along with the transducer cables, where they were protected in a pouch secured to the ewe’s flank. After recovery from anesthesia, the ewe was returned to the cage. Antibiotics (1 million units of penicillin G procaine and 100 mg of gentamicin sulfate) were administered intravenously to the ewe and into the amniotic cavity during surgery and daily thereafter. Buprenorphine (0.01 mg/kg im) was administered for postoperative analgesia. All protocols were approved by the Committee of Animal Research at the University of California, San Francisco.

Experimental protocol. After a 24-h recovery, the ewe was placed in a study cart with free access to food and water. The fetal catheters were connected to transducers, and 60 min were allowed for stabilization. PEG diluted in 5 ml of normal saline (n = 5, vehicle control) or polyethylene glycol-conjugated superoxide dismutase (n = 7, PEG-SOD) was then delivered through the pulmonary artery catheter. The dose of PEG-SOD (1,000–1,500 U/kg) was based on previous studies that demonstrate a sustained significant increase in plasma Mn-SOD (32, 33). The thoracotomy was then closed in layers. Warm saline was instilled to replace the lost amniotic fluid, and the uterine incision was closed. A polyvinyl catheter was placed in the amniotic cavity. The catheters were filled with PBS containing 1,000 U/ml heparin sodium, plugged, and brought to the skin along with the transducer cables, where they were protected in a pouch secured to the ewe’s flank. After recovery from anesthesia, the ewe was returned to the cage. Antibiotics (1 million units of penicillin G procaine and 100 mg of gentamicin sulfate) were administered intravenously to the ewe and into the amniotic cavity during surgery and daily thereafter. Buprenorphine (0.01 mg/kg im) was administered for postoperative analgesia. All protocols were approved by the Committee of Animal Research at the University of California, San Francisco.

In 12 of the fetal lambs, the vascular occluder placed around the ductus arteriosus was then inflated with normal saline, to increase mean pulmonary arterial pressure by 15–20 mmHg. The hemodynamic variables were monitored continuously, and systemic arterial blood gases were sampled intermittently. The occluder was occasionally adjusted to maintain the increase in mean pulmonary arterial pressure. This was required approximately once per animal, and there were no differences in the need for occluder manipulations between the two study groups. After 4 h, a repeat cesarean section was then performed, and a peripheral fetal lung biopsy was performed as described above.

To ensure that potential changes demonstrated resulted from ductal constriction, and not from other aspects of the protocol, six of the vehicle-treated fetal lambs underwent the exact protocol without inflation of the vascular occluder (sham operated).

At the end of the protocol, the fetus and ewe were killed with a lethal injection of pentobarbital sodium followed by bilateral thora-
maximum and minimum spectral amplitudes for the CM-superoxide spin-trap product waveform were quantified. Experimental groups were then compared for differences in amplitude using statistical analysis.

Preparation of protein extracts and Western blot analysis. Lung protein extracts were prepared by homogenizing peripheral lung tissues in Triton lysis buffer (50 mM Tris·HCl, pH 7.6, 0.5% Triton X-100, 20% glycerol) containing a protease inhibitor cocktail [P8340 (5 ml) was used to prevent protein degradation; P5726 (5 ml) was also used in the phospho-eNOS Western blots to prevent phosphate group cleavage, Sigma-Aldrich]. Extracts were then centrifuged (15,000 g × 10 min at 4°C). Supernatant fractions were then assayed for protein concentration using the Bradford reagent (Bio-Rad, Richmond, CA) and used for Western blot analysis. Western blot analysis was performed as previously described (4, 15, 30). Briefly, protein extracts (25 μg) were separated on 4–12% denaturing polyacrylamide gels and transferred to a nitrocellulose membrane (Bio-Rad). The membranes were blocked with 5% nonfat dry milk in TBS containing 0.1% Tween and exposed to the primary antibody. The primary antibodies used for immunoblotting were anti-phospho Thr495 eNOS, anti-phospho Ser1177 eNOS, and anti-eNOS (all at 1:1,000, Cell Signaling Technology). After blocking, the membranes were washed with TBS containing 0.1% Tween and incubated with the appropriate secondary antibody coupled to horseradish peroxidase, washed again with TBST, and the protein bands visualized with the appropriate secondary antibody coupling to horseradish peroxidase, washed again with TBST, and the protein bands visualized using chemiluminescence (SuperSignal West Femto Maximum Sensitivity Substrate Kit; Pierce, Rockford, IL) and a Kodak 440CF image station (New Haven, CT). To normalize for protein loading, blot were reblocked and then reprobed with the housekeeping protein, β-actin.

Immunoprecipitation-Western blot analysis for eNOS nitration. We determined the level of eNOS protein nitration utilizing an immunoprecipitation-Western blot technique as we have previously described (30). Frozen lung tissue was homogenized in 3X volume per tissue weight of immunoprecipitation (IP) buffer (25 mM HEPES, pH 7.5, 150 mM NaCl, 1% Nonidet P-40, 10 mM MgCl2, 1 mM EDTA, 2% glycerol supplemented with protease inhibitors), centrifuged at 14,000 rpm at 4°C for 10 min, the supernatant collected, and the protein concentration quantified by the Bio-Rad DC Protein Assay (Bio-Rad Laboratories, Hercules, CA). To 1,000 μg of total protein, 1 μg of anti-eNOS antibody was added, the volume was brought to 1 ml with immunoprecipitation buffer, and the mixture was incubated at 4°C overnight. To precipitate the bound eNOS, 10 μl of protein G-Agarose (EMD/Calbiochem) was added, and the samples incubated for 1 h at 4°C. To collect the bead-bound antibody, the samples were centrifuged at 14,000 rpm for 5 s, the supernatant was removed, and the beads were washed with 500 μl of IP buffer. The wash step was repeated two additional times, and 20 μl of 2X Laemmli sample buffer was added to the samples and boiled for 5 min. The samples were then divided equally and loaded onto duplicate 4–20% gradient gels (Gradiapore, Fresh荔枝 Forest, Australia) and run to completion according to the manufacturer’s instructions. The proteins were transferred to Immob-Blot polyvinylidene difluoride membrane (Bio-Rad Laboratories), and the membrane was blocked with 5% skim milk in TBST from 1 h to overnight. The membranes were probed with antibodies to either eNOS (to normalize for the immunoprecipitation efficiency) or 3-NT (EMD/Calbiochem), and reactive bands were determined as a ratio of the 3-NT-eNOS/total eNOS signals.

Table 1. Hemodynamic changes associated with acute ductal constriction in vehicle-treated fetal lambs

<table>
<thead>
<tr>
<th></th>
<th>Preconstriction</th>
<th>15 Min</th>
<th>30 Min</th>
<th>1 H</th>
<th>2 H</th>
<th>3 H</th>
<th>4 H</th>
</tr>
</thead>
<tbody>
<tr>
<td>PAP, mmHg</td>
<td>54.5 ± 3.7</td>
<td>66.6 ± 8.4*</td>
<td>65.2 ± 7.4*</td>
<td>68.8 ± 7.9*</td>
<td>68.2 ± 8.5*</td>
<td>64.7 ± 10.1*</td>
<td>65.5 ± 8.5*</td>
</tr>
<tr>
<td>Left pulmonary vascular resistance, mmHg</td>
<td>20.3 ± 19.6</td>
<td>3.3 ± 2.5</td>
<td>2.3 ± 1.2*</td>
<td>4.3 ± 2.7</td>
<td>16.3 ± 15.2</td>
<td>20.7 ± 16.6*</td>
<td>35.7 ± 29.9†</td>
</tr>
<tr>
<td>Left pulmonary blood flow, ml kg⁻¹ min⁻¹</td>
<td>7.3 ± 6.6</td>
<td>21.1 ± 9.9*</td>
<td>25.9 ± 10.9*</td>
<td>17.7 ± 12.4</td>
<td>8.1 ± 8.0†</td>
<td>4.4 ± 4.4†</td>
<td>3.8 ± 5.4†</td>
</tr>
<tr>
<td>SAP, mmHg</td>
<td>53.0 ± 2.6</td>
<td>49.5 ± 6.1</td>
<td>50.9 ± 4.3</td>
<td>49.8 ± 4.9</td>
<td>49.1 ± 7.5</td>
<td>46.1 ± 10.3</td>
<td>48.0 ± 6.2</td>
</tr>
<tr>
<td>SAP-PAP, mmHg</td>
<td>1.4 ± 1.6</td>
<td>17.1 ± 3.7*</td>
<td>16.1 ± 4.5*</td>
<td>19.0 ± 6.1*</td>
<td>19.2 ± 5.1*</td>
<td>18.5 ± 8.3*</td>
<td>17.5 ± 4.6*</td>
</tr>
<tr>
<td>PH, units</td>
<td>7.31 ± 0.16</td>
<td>7.42 ± 0.07</td>
<td>7.40 ± 0.07</td>
<td>7.41 ± 0.08</td>
<td>7.37 ± 0.05</td>
<td>7.37 ± 0.08</td>
<td>7.38 ± 0.09</td>
</tr>
<tr>
<td>PaO₂, Torr</td>
<td>47.4 ± 9.4</td>
<td>43.0 ± 5.9</td>
<td>42.7 ± 8.2</td>
<td>42.7 ± 8.2</td>
<td>44.4 ± 6.8</td>
<td>45.4 ± 8.8</td>
<td>47.6 ± 9.9</td>
</tr>
<tr>
<td>PaCO₂, Torr</td>
<td>16.7 ± 3.1</td>
<td>16.0 ± 2.0</td>
<td>16.3 ± 2.5</td>
<td>17 ± 1.7</td>
<td>15.7 ± 3.1</td>
<td>16 ± 4.5</td>
<td>15.5 ± 2.6</td>
</tr>
</tbody>
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Values are means and SD; n = 5 lambs. PAP, pulmonary arterial pressure; SAP, systemic arterial pressure. *P < 0.05 vs. preconstriction values. †P < 0.05 vs. previous column.

Fig. 1. Changes in left pulmonary blood flow (LPA; A) and left pulmonary vascular resistance (LPVR; B) before and after acute ductal constriction. Acute ductal constriction induces an initial increase in pulmonary blood flow and decrease in pulmonary vascular resistance. This is followed by active pulmonary vasoconstriction, resulting in a decrease in flow and increase in resistance. Pretreatment with PEG-SOD attenuated this response. N = 5 vehicle-treated lambs, n = 7 PEG-SOD-treated lambs. Values are means ± SD. *P < 0.05 vs. pre, †P < 0.05 vs. 30 min, ‡P < 0.05 between groups.
Statistical analysis. Means ± SD were calculated for the protein of interest. The means ± SD were calculated for the hemodynamic variables, NOS activities, and EPR amplitudes. Differences over time were determined by ANOVA for repeated measures, and Student-Newman-Keuls post hoc testing was performed. Values between groups were compared by ANOVA for repeated measures. A P value of less than 0.05 was considered statistically significant.

RESULTS

There were no differences in gestational age, weight, sex distribution, or baseline hemodynamic variables between vehicle-treated, PEG-SOD-treated, and non-constricted fetal lambs (data not shown).

In vehicle (PEG-alone)-treated lambs, acute ductal constriction rapidly increased mean pulmonary arterial pressure and left pulmonary blood flow (P < 0.05). Left pulmonary vascular resistance decreased (P < 0.05). Mean systemic arterial pressure, left atrial pressure, and systemic arterial blood gases and pH were all unchanged (Table 1). During the 4-h study period, pulmonary arterial pressure remained increased (as per protocol), but left pulmonary blood flow and pulmonary vascular resistance returned to preconstriction values (Fig. 1). In fact, compared with 30-min postconstriction, pulmonary blood flow was decreased after 4 h, and pulmonary vascular resistance was increased (P < 0.05, Fig. 1).

In vehicle-treated lambs, NOS activity decreased by ~50% (P < 0.05, Fig. 2) 4 h following acute ductal constriction. These changes were independent of changes in eNOS protein levels and are consistent with our previous findings (17). Superoxide levels, as determined by EPR on peripheral lung, increased following acute ductal constriction in vehicle-treated fetal lambs (P < 0.05, Fig. 3). Specificity of the EPR assay for superoxide was confirmed by a significant reduction in the waveform amplitude with the addition of PEG-SOD to the samples.

Posttranslational alterations in NOS can result in changes in NOS activity. Since ROS can alter both the nitration and phosphorylation status of NOS, we evaluated potential alterations in nitroated eNOS, Ser1177 eNOS, and eNOS Thr495 following acute constriction of the ductus arteriosus. As seen in Fig. 4, total amounts of nitroated eNOS and Ser1177 eNOS were not changed following acute ductal constriction. However, the amount of eNOS Thr495, which is associated with decreased NOS activity, was increased following acute ductal constriction (P < 0.05).

To determine potential NO-ROS interactions following ductal constriction, an additional group of fetal lambs was pretreated with PEG-SOD. In PEG-SOD-treated lambs, acute ductal constriction rapidly increased mean pulmonary arterial pressure and left pulmonary blood flow (P < 0.05). Left pulmonary vascular resistance decreased (P < 0.05). Mean systemic arterial pressure, left atrial pressure, and systemic arterial blood gases and pH were all unchanged (Table 2). During the 4-h study period, pulmonary arterial pressure remained increased (as per protocol), but left pulmonary vascular resistance remained unchanged (Table 2, Fig. 1). Left pulmonary blood flow decreased at hour 3 and 4 but remained higher than preconstriction values. In fact, after 3 and 4 h of ductal constriction, pulmonary blood flow was significantly increased, and pulmonary vascular resistance was significantly decreased in PEG-SOD-treated lambs compared with vehicle-treated lambs (P < 0.05, Fig. 1).

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In PEG-SOD-treated lambs, lung tissue NOS activity did not decrease following acute ductal constriction, although eNOS protein levels were unchanged (Fig. 2). Superoxide levels and 3-NT eNOS levels were also unchanged (Figs. 3 and 4). Interestingly, although phospho-Thr495 eNOS levels were increased as in vehicle-treated lambs, levels of phospho-Ser1177 eNOS were not affected by PEG-SOD treatment. In fact, when compared side by side, phospho-Ser1177 eNOS protein levels were greater in the PEG-SOD-treated lambs following ductal constriction than the PEG-alone-treated lambs (G). N = 5 vehicle-treated lambs, n = 7 PEG-SOD-treated lambs. Values are means ± SD. *P < 0.05 vs. Pre.

No changes in hemodynamic variables, superoxide levels, NOS activity, and protein levels were noted in non-constricted (sham) fetal lambs (data not shown).

DISCUSSION

The regulation of the high basal pulmonary vascular tone, and its dramatic transition to a low-resistance circulation after birth, involves a complex, incompletely understood, interaction between anatomic alterations, mechanical forces, and a balance between varieties of vasoactive mediators with competing effects on basal tone. Although the maintenance of low fetal
pulmonary blood flow and high pulmonary vascular resistance is considered adaptive given the placental circulation and the resulting non-dependence of pulmonary blood flow for gas exchange, aberrations in these mechanisms may result in incomplete transition to the low postnatal pulmonary resistance and persistent pulmonary hypertension of the newborn (PPHN). Recent studies have demonstrated a role for increased myogenic tone in the maintenance of the high fetal pulmonary vascular resistance, but their mechanisms are incompletely understood. Previously, we have demonstrated a role for decreased eNOS activity in the intense pulmonary vasoconstriction following an increase in pulmonary blood flow secondary to acute ductal constriction in the fetal lamb (17). In the current study, we found that the decrease in eNOS activity is associated with an increase in lung superoxide levels and changes in the phosphorylation status of eNOS. Furthermore, we found that pretreatment with PEG-SOD attenuates the vasoconstriction and preserves NOS activity, suggesting a role for ROS-eNOS interactions in the pulmonary myogenic response following acute ductal constriction in the fetus.

A novel finding of this study is the demonstration of an 82% increase in lung superoxide levels 4 h following acute constriction of the fetal ductus arteriosus (Fig. 3). Several potential mechanisms may be involved in this finding, including alterations in mechanical forces, ET-1, and the phosphorylation status of eNOS. Ductal constriction results in a sustained increase in pulmonary arterial pressure and a transient increase in flow. Interestingly, in vitro studies demonstrate that the mechanical forces associated with increased pressure (cyclic mechanical strain) increase superoxide production via an up-regulation of NADPH oxidase (11, 14, 28), suggesting a possible mechanism for the increase in superoxide demonstrated in this study. In addition, we have previously shown that ET-1 levels are increased following acute ductal constriction, and in vitro data demonstrate that ET-1 can increase superoxide production in isolated fetal pulmonary artery smooth muscle cells (29). Last, the current study demonstrates changes in the phosphorylation status of eNOS, and data suggest that such changes may uncouple eNOS resulting in NOS-derived superoxide production (9). The exact mechanisms involved in the increase in superoxide production in this study warrant further investigation.

Associated with the increase in superoxide reduction was a decrease in eNOS activity in the intense pulmonary vasoconstriction following an acute constriction of the fetal ductus arteriosus (Fig. 3). Several posttranslational factors, including the nitrational and phosphorylation status of the enzyme, may dynamically regulate eNOS activity (10, 18, 20, 21), and, interestingly, several lines of evidence suggest that ROS can modulate many of these posttranslational alterations. For example, superoxide reacts rapidly with NO to form peroxynitrite, a strong oxidizing agent, which reacts readily with biological molecules and is capable of nitrating free or protein-associated tyrosines, including eNOS, rendering it inactive. In the current study, we found no differences in the amounts of nitrated eNOS following 4 h of acute ductal constriction between the PEG-alone- and PEG-SOD-treated lungs, suggesting that eNOS nitration was not responsible for the differences in eNOS activity between the two groups (Fig. 4). However, because of limitations in the amount of preconstriction tissue available, comparisons between pre- and postconstriction in the PEG-alone tissues could not be assessed to completely rule out the responsibility of changes in eNOS nitration in this preparation. Phosphorylation and dephosphorylation of eNOS is another important posttranslational modification that regulates activity. Although there are numerous phosphorylation sites, the best studied are the phosphorylation sites Thr495 and Ser1177. Phosphorylation of Ser1177 induces eNOS activation, whereas phosphorylation of Thr495 is inhibitory (3). We found that acute compression of the fetal ductus arteriosus resulted in a significant increase in eNOS-Thr495, whereas eNOS-Ser1177 was unchanged. The net result of these changes could explain, at least in part, the associated decrease in NOS activity. Other posttranslational modifications, such as intracellular location, protein-protein interactions, and substrate and cofactor availability, could participate in the changes in NOS activity during ductal constriction, and further investigations are planned to elucidate these complex processes.

Pretreatment with PEG-SOD prevented the reduction in NOS activity and attenuated the intense pulmonary vasoconstriction following ductal constriction, suggesting a role for ROS in this myogenic response. This lack of a decrease in NOS activity induced by PEG-SOD treatment was not associated with a decrease in eNOS-Thr495 levels, a change in nitrated eNOS, or a change in eNOS protein expression. Interestingly, PEG-SOD treatment was associated with an increase in the lung protein levels of the stimulatory eNOS-
Ser1177, suggesting that the increase in eNOS-Ser1177, at least in part, is responsible for the restoration in NOS activity. However, ROS can be involved in several other potential NOS modifications that require further investigation.

Several important limitations of the present study are noteworthy. First, only one ROS, superoxide, was studied. Other ROS, such as hydrogen peroxide, may be altered by mechanical forces and may alter NOS activity, and thus warrant further study (34). Second, the model utilized in this study employs peripheral lung biopsies. Therefore, the biochemical analysis is performed on peripheral lung homogenate, excluding the possibility of examining specific arterial sites of the myogenic response within the lung vasculature, and isolating changes to specific cell types. Strengths of this study include the in vivo performance of these investigations in chronically instrumented late-gestation fetal sheep, and the preconstriction lung biopsy allows the determination of changes within the same animal over time.

We conclude that alterations in ROS participate in the hemodynamic and biochemical alterations associated with the myogenic response following fetal constriction of the ductus arteriosus. We speculate that aberrations in these adaptive mechanisms may participate in the pathophysiology of PPHN and the alterations in fetal blood flow patterns associated with certain congenital heart defects, and thus warrant further investigation.

GRANTS

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DISCLOSURES

No conflicts of interest (financial or otherwise) are declared by the author(s).

REFERENCES


