TRANSLATIONAL PHYSIOLOGY

Therapeutic potential of sustained-release sodium nitrite for critical limb ischemia in the setting of metabolic syndrome

David J. Polhemus,1* Jessica M. Bradley,1 Kazi N. Islam,1 Luke P. Brewster,2,3,4 John W. Calvert,2
Ya-Xiong Tao,5 Carlos C. Chang,6 Iraklis I. Pipinos,7 Traci T. Goodchild,1 and David J. Lefer1

1Cardiovascular Center of Excellence and Department of Pharmacology, LSU Health Sciences Center, New Orleans, Louisiana; 2Department of Surgery, Emory University School of Medicine, Atlanta, Georgia; 3Surgery and Research Services, Atlanta Veterans Affairs Medical Center, Decatur, Georgia; 4Parker H. Petit Institute for Bioengineering and Bioscience, Georgia Institute of Technology, Atlanta, Georgia; 5Department of Anatomy, Physiology, and Pharmacology, Auburn University College of Veterinary Medicine, Auburn, Alabama; 6T3 Labs, Emory University, Atlanta, Georgia; and 7Department of Cellular and Integrative Physiology, University of Nebraska Medical Center, Omaha, Nebraska

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Polhemus DJ, Bradley JM, Islam KN, Brewster LP, Calvert JW, Tao YX, Chang CC, Pipinos II, Goodchild TT, Lefer DJ. Therapeutic potential of sustained-release sodium nitrite for critical limb ischemia in the setting of metabolic syndrome. Am J Physiol Heart Circ Physiol 309: H82–H92, 2015. First published April 24, 2015; doi:10.1152/ajpheart.00115.2015.—Nitrite is a storage reservoir of nitric oxide that is readily reduced to nitric oxide under pathological conditions. Previous studies have demonstrated that nitrite levels are significantly reduced in cardiovascular disease states, including peripheral vascular disease. We investigated the cytoprotective and proangiogenic actions of a novel, sustained-release formulation of nitrite (SR-nitrite) in a clinically relevant in vivo swine model of critical limb ischemia (CLI) involving central obesity and metabolic syndrome. CLI was induced in obese Ossabaw swine (n = 18) by unilateral external iliac artery deployment of a full cross-sectional vessel occlusion device positioned within an endovascular expanded polytetrafluoroethylene-lined nitinol stent-graft. At post-CLI day 14, pigs were randomized to placebo (n = 9) or SR-nitrite (80 mg, n = 9) twice daily by mouth for 21 days. SR-nitrite therapy increased nitrate, nitrite, and S-nitrosothiols in plasma and ischemic skeletal muscle. Oxidative stress was reduced in ischemic limb tissue of SR-nitrite- compared with placebo-treated pigs. Ischemic limb tissue levels of proangiogenic growth factors were increased following SR-nitrite therapy compared with placebo. Despite the increases in cytoprotective and angiogenic signals with SR-nitrite therapy, new arterial vessel formation and enhancement of blood flow to the ischemic limb were not different from placebo. Our data clearly demonstrate cytoprotective and proangiogenic signaling in ischemic tissues following SR-nitrite therapy in a very severe model of CLI. Further studies evaluating longer-duration nitrite therapy and/or additional nitrite dosing strategies are warranted to more fully evaluate the therapeutic potential of nitrite therapy in peripheral vascular disease.

nitric oxide; metabolic syndrome; angiogenesis; peripheral arterial disease; Ossabaw swine; endothelial nitric oxide synthase; nitrite
of NO that is readily reduced to NO under pathological conditions including ischemia (15, 30). Nitrite has been shown to be cytoprotective in numerous animal models of ischemic injury, including myocardial ischemia-reperfusion injury (19, 31, 38), heart failure (6), cerebral vasospasm (36), and hindlimb ischemia (6, 10, 19, 26). In a recent phase II PAD clinical trial (i.e., SONIC), sodium nitrite therapy was shown to be safe and well tolerated (32). A major limitation for the clinical trial (i.e., SONIC), sodium nitrite therapy was shown to be effective in the treatment of CLI and MetS.

METHODS

Human CLI and control skeletal muscle collection. After removal of the affected extremity, the amputation specimen was transferred to a separate sterile table, and elliptical 8 cm muscle biopsies of the anterior tibialis and gastrocnemius muscles were performed. For above-knee amputations, an additional biopsy of the vastus medialis was performed. All patients consented to the use of their amputated tissue for research purposes and signed a Health Insurance Portability and Accountability Act release for use of their medical records. Patients were segregated into normal perfusion or CLI according to the presence of a Doppler signal. All patients were included in the CLI group.

Animals. Castrated male Ossabaw swine, 6–8 mo of age, were obtained from Indiana University. Although female swine display a more rapid development of dyslipidemia and atherosclerosis when fed an atherogenic obese diet (1 kg/day) consisting of 2% cholesterol, 17% coconut oil, 2.5% corn oil, and 0.7% sodium cholate (KT324, Purina Test Diet).

Sodium nitrite therapy. SR-nitrite tablets were generously supplied by TheraVasc (Cleveland, OH). Animals received placebo (n = 9) or SR-nitrite (80 mg, n = 9) by mouth twice daily. Treatment was initiated at 14 days following induction of CLI and continued for 21 days (Fig. 1). CLI model. Obese male swine (n = 18) were subjected to CLI as depicted in Fig. 1. Nine pigs were originally assigned to the placebo group and nine to the SR-nitrite group in a blinded and randomized fashion. During the course of the 35-day CLI protocol, three pigs in the placebo group and two in SR-nitrite group died as a result of complications of CLI and obesity/MetS. Therefore, there were six pigs in the placebo group and seven in the SR-nitrite group at day 35. Animals were sedated with ketamine (15 mg/kg) in combination with xylazine (1 mg/kg) and, to aid induction, were endotracheally intubated and maintained on isoflurane in oxygen. The preoperative analgesic buprenorphine (0.15 mg for each pig) was administered along with the perioperative antibiotic cefazolin (1,000 mg). Postoperative analgesics, carprofen (10 mg) and buprenorphine (0.15–0.3 mg), were given immediately after the procedure. On postoperative day 1, additional analgesics, buprenorphine (0.3 mg) in combination with carprofen (100 mg), were administered. Heart rate, respiration, peripheral capillary O2 saturation, end-tidal CO2, fluid volume, and body temperature were monitored throughout the procedure. Activated clotting time was measured with a coagulation-monitoring system (Hemachron) after placement of the arterial sheath and administration of heparin.

Under sterile conditions and with fluoroscopic guidance, an 8-French introducer (Maximum Hemostasis Introducer 23-cm ACT Sheath, St. Jude Medical, St. Paul, MN) was inserted into the right external carotid artery, advanced into the abdominal aorta, and placed just above the aortoiliac bifurcation. Heparin (300 U/kg) was administered, and activated clotting time was monitored throughout the procedure to ensure clotting times of ≥250 s. A percutaneous guide wire (0.035 × 230 cm; Rosen Starter, Boston Scientific, Natick, MA) was positioned in the right external iliac artery. A 7-French delivery guide catheter (Mach 1 MP1, Boston Scientific) containing a self-expanding endoluminal endoprosthesis consisting of an expanded polytetrafluoroethylene (ePTFE) lining with an external nitinol stent (6 mm × 15 cm; VIABAHN, W. L. Gore and Associates, Flagstaff, AZ) was advanced, positioned within the right external iliac artery, and deployed. Contrast arteriography was performed to evaluate vessel patency. After placement of the ePTFE-lined endoprosthesis, a self-expanding nitinol mesh occlusion device (8 × 7 mm; AMPLATZER Vascular Plug II, St. Jude Medical) was loaded onto the delivery guide catheter, advanced, positioned within the right external iliac artery, and deployed within the proximal portion of the ePTFE-lined stent.

![Critical Limb Ischemia in Obese Ossabaw Swine](image)

Fig. 1. Critical limb ischemia (CLI) protocol. Obese Ossabaw swine were subjected to 35 days of CLI. Sustained-release (SR)-nitrite therapy was initiated at a dose of 80 mg by mouth twice a day beginning 14 days after induction of CLI. Vascular angiographic images were obtained at baseline, immediately post-CLI, and on day 35. Animals were euthanized after 35 days of CLI, and skeletal muscle was collected for molecular and pharmacokinetic analyses. RSNO, S-nitrosothiol.
An arteriogram was performed to verify occlusion. The guide wire, guide catheter, and introducer sheath were removed, and the left external carotid artery was ligated. The incision was closed, the animal was allowed to recover, and buprenorphine (0.05 mg/kg) was administered for analgesia. The ePTFE-lined endoprosthesis served to inhibit recanalization of the artery by collateral vessels, typically observed in animal models of CLI.

Euthanasia and tissue collection. After all in vivo measurements were completed, heparin (200 U/kg iv) was injected and allowed to circulate for ~5 min. Animals were euthanized while under deep inhalant anesthesia (isoflurane, 5%). The gastrocnemius muscle was collected from control and CLI limbs.

Measurement of NO metabolites. Nitrite, nitrate, S-nitrosothiol (RSNO), and NO-heme concentrations were quantified as previously described (6). Immediately after euthanasia, ischemic and control skeletal muscles were removed and flash-frozen using liquid nitrogen. Tissue was homogenized in PBS (10 ml/mg of tissue). In some cases, differences in sample size are due to a limited quantity of adequate tissue samples required for these complex assays.

Angiography. Hindlimb vasculature was visualized during record-ings using undiluted contrast (Oxilan 300, Guerbet, Bloomington, IN). Arteriographs of the uninstrumented left and occluded right iliac arteries were obtained at baseline and on day 35.

Ankle-brachial index. An automated blood pressure machine with a 2-inch cuff was used to measure limb blood pressure before CLI, immediately post-CLI, and at the end of the experiment. Animals were sedated using isoflurane in oxygen by mask to effect, and a pressure cuff was secured to each limb, rapidly inflated to ~30 mmHg above the anticipated systolic pressure, and then deflated slowly for determination of systolic blood pressure in each of the forelimbs and hindlimbs. Ankle-brachial index (ABI) was calculated for each side of the animal by dividing the systolic blood pressure measured at the hindlimb (ankle) by that measured at the contralateral hindlimb. In some cases, we were unable to obtain reproducible or reliable cuff readings for ABI data; in these cases, ABI was not reported. Therefore, the sample size reported for the ABI data is variable.

cGMP RIA. Skeletal muscle cGMP concentrations were quantified as previously described (6).

Western blot analysis. Gastrocnemius muscle was used for Western blot analysis, performed as described previously (25). The following primary antibodies were used: VEGF (Abcam), CD31 (Novus Biologicals), Ser1177-phosphorylated endothelial NO synthase (eNOS; Cell Signaling), and eNOS, Thr495-phosphorylated eNOS, and GAPDH (Santa Cruz Biotechnology).

Real-time PCR. Skeletal muscle mRNA was quantified using Taq-Man primers (Life Technologies) as previously described (34).

Quantitative angiographic score. Collateral vessel growth in the control and CLI limbs was assessed using a grid overlay of 2-mm squares (44). Quantitative vessel analysis was performed on angiographic images of the control and CLI limbs distal from the ePTFE-lined endoprosthesis at post-CLI day 35. The angiogenic score was calculated as percentage of the number of contrast-opacified vessels crossing the squares divided by the total number of vessels. Sample size was limited to five per group because of technical complications during angiographic imaging as well as the high mortality rate in the Ossabaw swine subjected to CLI.

ELISAs. VEGF, CD31, and von Willebrand factor were assayed using ELISA kits according to the manufacturer’s recommendations (MyBioSource, San Diego, CA).

Determination of protein carbonyl content. Protein carbonyl content of skeletal muscle was measured as described previously (25).

Measurement of malondialdehyde levels. Malondialdehyde (MDA) levels were measured in skeletal muscle as previously described (25).

Statistical analysis. Values are means ± SE. Differences in data between the groups were compared using Prism 4 (GraphPad Software) with Student’s unpaired, two-tailed t-test when only two groups were compared. P < 0.05 was considered statistically significant.

RESULTS

CLI patients exhibit NO deficiency. The NO intermediates (nitrite, nitrosothiols, and NO-heme), as well as cGMP and cGMP-dependent protein kinase-1 (PKG-1), were measured in gastrocnemius, anterior tibialis, and vastus medialis tissue obtained postamputation from normal and CLI patients (Fig. 2). Patient data for the control and CLI patients are presented in Table 1. We observed significant (P < 0.01) reductions in skeletal muscle nitrite (Fig. 2A), RSNO (Fig. 2B), and NO-heme (Fig. 2C) levels in CLI patients compared with age-matched controls. Skeletal muscle cGMP levels were also significantly (P < 0.01) attenuated in CLI patients compared with controls (Fig. 2D). Furthermore, PKG-1 protein expression and mRNA levels were significantly (P < 0.01) reduced in CLI tissues compared with controls (Fig. 2, G and H).

Obese Ossabaw swine exhibit eNOS dysfunction and are NO-deficient. Western blot analysis of soleus muscle obtained from lean and obese swine revealed no change in total eNOS protein expression (Fig. 3B). However, there was a significant increase in phosphorylation at the eNOS inhibitory site, Thr495, in the obese swine (Fig. 3C). There was no difference in phosphorylation status at the eNOS activation site, Ser1177, between groups (Fig. 3D). Asymmetric dimethylarginine (ADMA) is structurally similar to L-arginine and inhibits NO synthase, resulting in decreased NO production and endothelial dysfunction. The obese swine displayed elevated circulating ADMA levels compared with the lean group (Fig. 3E). Nitrite, a marker for NO bioavailability, was significantly (P < 0.05) decreased in plasma and skeletal muscle of the obese animals compared with the lean controls (Fig. 3F), providing further evidence of NO insufficiency in the obese Ossabaw animals.

Pharmacokinetics of SR-nitrite in CLI. Beginning at post-CLI day 14, SR-nitrite (80 mg) or placebo was administered by mouth twice daily for 21 days (Fig. 4A). We did not observe a significant change in plasma nitrite or nitrate in the placebo-treated group during the 35-day CLI experimental protocol. In contrast, we observed an increase in circulating nitrite levels following SR-nitrite therapy, with a statistically significant (P < 0.05) difference between the SR-nitrite- and placebo-treated groups at day 35. We failed to observe statistically significant increases in circulating nitrite levels at days 21 and 28. This is possibly due to immediate reduction of nitrite to NO and NO scavenging in the circulation due to severe ischemia and oxidative stress in the setting of CLI. Additionally, plasma nitrate levels were markedly increased following SR-nitrite administration at days 21 (P < 0.01), 28 (P < 0.01), and 35 (P < 0.05; Fig. 2C). Furthermore, we observed an increase in skeletal muscle tissue nitrite (Fig. 4B) and nitrate (Fig. 4D) levels following SR-nitrite therapy, as evidenced by a fourfold increase (P < 0.01) in nitrite in the CLI limb compared with the placebo-treated group. We also examined RSNO concentrations in plasma and skeletal muscle. At day 35, plasma RSNO levels were significantly (P < 0.05) greater in the nitrite- than placebo-treated group. We observed a 3.9-fold increase in tissue RSNO concentration in the CLI limb of the nitrite- compared with placebo-treated swine (Fig. 4E). These data indicate that nitrite levels are increased in the ischemic limb and result in increased NO bioavailability. Safety of augmented nitrite levels following therapy was validated by stable circulating methemoglobin levels in the SR-nitrite-
treated group at day 35. There was no difference in percent
methemoglobin between the SR-nitrite- and placebo-treated
groups (3.1 ± 0.9% vs. 2.1 ± 0.5%, P = NS).

Nitrite therapy fails to improve extremity blood flow in CLI.
Measurement of blood flow revealed an immediate reduction in
ABI to 0 following the CLI procedure. This reduction gradu-
ally improved during the 35-day CLI protocol but continued to
demonstrate a persistent ischemic state. ABI (Fig. 5) revealed
no significant change in blood flow of the ischemic limb
comparing the SR-nitrite- vs. placebo-treated group.

Contrast angiography. Representative, magnified-view con-
trast angiographic images of Ossabaw swine peripheral vascu-
lature obtained at baseline, immediately after induction of CLI,
and at post-CLI day 35 are shown in Fig. 6A.

Hindlimb vessel density was determined from the angio-
graphic score of the magnified-view contrast angiographic
images for the control and CLI limb distal from the ePTFE-
lined endoprosthesis at post-CLI day 35. Quantitative angio-
graphic analysis (Fig. 6B) revealed a significant reduction in
perfusion of the CLI limb (P < 0.01) in the placebo-treated
animals between the control and CLI limb at post-CLI day 35.

Angiographic assessment of hindlimb perfusion at day 35
revealed no significant changes in the CLI limb of SR-nitrite-
treated animals compared with the placebo-treated group.

Nitrite therapy promotes angiogenic signaling in the isch-
emic limb. Quantitative PCR and Western blot analysis per-
formed at post-CLI day 35 demonstrate increases in known
angiogenic signals following SR-nitrite therapy (Fig. 7). VEGF

Table 1. Patient information for control and CLI patients
who underwent extremity amputations

<table>
<thead>
<tr>
<th>Group</th>
<th>Control (n = 4)</th>
<th>CLI (n = 16)</th>
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<tbody>
<tr>
<td>Age, yr</td>
<td>64.5 ± 13.44</td>
<td>68.1 ± 14.24</td>
</tr>
<tr>
<td>Male, %</td>
<td>0</td>
<td>63</td>
</tr>
<tr>
<td>Diabetes, %</td>
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<td>50</td>
</tr>
<tr>
<td>Hypertension, %</td>
<td>50</td>
<td>63</td>
</tr>
<tr>
<td>Congestive heart failure, %</td>
<td>0</td>
<td>25</td>
</tr>
<tr>
<td>Current/former smoker, %</td>
<td>0</td>
<td>31</td>
</tr>
</tbody>
</table>

Values for age are means ± SD. Skeletal muscle biopsies were obtained from anterior tibialis and gastrocnemius muscles. CLI, critical limb ischemia.

Fig. 2. CLI patients exhibit nitric oxide (NO) deficiency
in the ischemic limb. A–E: nitrite, RSNO, NO-heme,
cGMP, and nitrate in gastrocnemius muscle from CLI
patients and controls. F: representative Western blot of
gastrocnemius cGMP-dependent protein kinase 1 (PKG-
1). G and H: PKG-1 RNA message and PKG-1 protein
expression. Numbers inside circles denote number of
patients.
mRNA (Fig. 7A) was increased by approximately twofold in the ischemic limb following SR-nitrite treatment ($P < 0.05$ vs. placebo). Platelet endothelial cell adhesion molecule-1 (or CD31) is a marker for endothelial cell expansion, and we did not observe an increase in CD31 mRNA in the CLI limb of SR-nitrite-treated compared with placebo-treated animals ($P = 0.51$; Fig. 7B). Representative Western blots for VEGF and CD31 are shown in Fig. 7C. VEGF protein expression trended higher ($P = 0.051$) in the SR-nitrite- than placebo-treated CLI limb, while SR-nitrite failed to alter VEGF protein expression in the control limb (Fig. 7D). CD31 protein expression was significantly ($P < 0.05$) increased in the CLI limb of the SR-nitrite-group compared with the placebo-treated group (Fig. 7E). Skeletal muscle protein levels of basic fibroblast growth factor (Fig. 7F) were unchanged in control and CLI limbs of placebo- and SR-nitrite-treated animals. However, we did observe a significant ($P < 0.05$) increase in skeletal muscle von Willebrand factor in the CLI limb of the SR-nitrite-treated group (Fig. 7G). Collectively, these data suggest that SR-nitrite therapy did result in increased angiogenic activity in ischemic hindlimb skeletal muscle.

**SR-nitrite attenuates skeletal muscle oxidative stress and activates eNOS following CLI.** One of the pathological consequences of chronic tissue ischemia is increased oxidative stress and cellular injury. Since NO is known to modulate redox status and mitigate oxidative stress, we evaluated the extent of oxidative stress in control and CLI skeletal muscle at post-CLI day 35. Skeletal muscle levels of carbonyl and MDA are shown in Fig. 8. Carbonyl levels (Fig. 8A) were unchanged in the control limb following CLI. However, carbonyl levels were significantly ($P < 0.05$) elevated in the CLI limb compared with the control limb in placebo-treated animals. Treatment with SR-nitrite significantly ($P < 0.05$) attenuated carbonyl levels in the CLI limb compared with placebo treatment. Similar results for MDA are shown in Fig. 8B. No changes in MDA levels were observed in the control limb following CLI. MDA levels were significantly ($P < 0.05$ vs. control) increased in the CLI limb of placebo-treated animals compared with the control limb. SR-nitrite failed to significantly attenuate MDA levels in the CLI limb compared with placebo treatment. These studies confirm previous findings that nitrite and NO significantly attenuate oxidative stress during pathological states (23, 42). Recent studies have shown that nitrite therapy activates eNOS and that a reduction in oxidative stress also recouples eNOS to its more functional dimerized form (6, 11, 25). We observed that SR-nitrite therapy resulted in improved eNOS functionality in the ischemic skeletal muscle, as evidenced by increased Ser$^{1177}$ phosphorylation of eNOS compared with placebo treatment (Fig. 8, C and E). There was no change in total eNOS expression between groups (Fig. 8, C and D).

**DISCUSSION**

Effective new treatments for CLI are critically important, given the high incidence of major amputation and mortality in
this population (22). Increasing blood flow, tissue perfusion, and oxygen and nutrient delivery via new vessel growth or collateralization could promote tissue regeneration and/or delay or prevent tissue necrosis and the development of conditions such as gangrene. Since ~50% of CLI patients are poor candidates for revascularization because of unavailable anatomy or conduit (37), we sought a pharmacological approach to enhance perfusion, attenuate skeletal muscle injury, and reduce necrosis in a severe model of CLI in the setting of obesity and MetS. In the present study we evaluated the effects of delayed therapy with a novel, SR-nitrite formulation on severe CLI in a clinically relevant porcine model of obesity and MetS. Our data clearly indicate that oral administration of SR-nitrite is well tolerated, increases circulating and ischemic zone tissue levels of nitrite and other NO intermediates, reduces oxidative stress in ischemic tissue, and triggers proangiogenic and cytoprotective tissue signaling in the setting of severe CLI. However, we failed to observe significant increases in ischemic zone blood flow or vascularity following SR-nitrite treatment in CLI.

Previous studies of limb ischemia have been primarily limited to murine and rabbit model systems, in which CLI is introduced via surgical ligation of the femoral artery (16, 40). This procedure often results in substantial angiogenic and arteriogenic responses, with complete restoration of hindlimb blood flow in a relatively short time frame. This response to
A surgical occlusion of a peripheral artery is inconsistent with the pathophysiology of PAD or CLI in humans (12, 26, 43). Therefore, we sought to create a more clinically relevant model of CLI in the setting of obesity/MetS. Angiographic analysis and ABI measurements immediately following CLI induction clearly demonstrate complete occlusion of the external iliac artery and undetectable ankle blood pressure. At post-CLI day 35, angiography revealed a nearly complete absence of significant collateral vessel growth at the location of the stent.

It is well established that blood vessels require continuous production of NO (29), and reductions in NO bioavailability significantly impact vascular function, resulting in vasoconstriction, thrombosis, atherogenesis, and hypertension (17). Initial experiments demonstrated significant reductions in NO bioavailability and NO-mediated signaling in human gastrocnemius muscle of CLI tissue compared with healthy control tissue. Böger et al. (8) observed a correlation between gradual reduction of NO synthesis and increased symptoms in PAD patients. NO deficiency, likely due to the combination of NO scavenging and NO synthase dysregulation, results in profound reductions in NO bioavailability and vascular dysfunction. In addition to being a potent vasodilator, antiapoptotic agent, and antioxidant, NO has also been shown to mediate angiogenesis and promote blood flow in response to tissue ischemia (14, 33).

Fig. 6. Contrast angiographic images and angiographic score at post-CLI day 35. A: magnified-view contrast angiographic images of swine peripheral vasculature at baseline, immediately post-CLI, and at day 35 post-CLI after interventional deployment of a self-expanding endoluminal endoprosthesis consisting of an expanded polytetrafluoroethylene (ePTFE) lining with an external nitinol stent (black arrow) and a self-expanding nitinol mesh occlusion device (white arrow) into the right external iliac artery. B: hindlimb vessel density shown as angiographic score of magnified-view contrast angiographic images of control and CLI limbs distal from the ePTFE-lined endoprosthesis at CLI day 35. Numbers inside circles denote number of animals. **P < 0.01 vs. placebo-treated control limb.

Control Limb | CLI Limb
--- | ---
Placebo | SR-Nitrite

Left axis: Placebo | Right axis: SR-Nitrite

### Variables
- Angiographic Score (%)
- Placebo
- SR-Nitrite

P = NS

10 15 20

Day 35 Baseline Immediately Post CLI
Specifically, NO therapy with sodium nitrite increases ischemic hindlimb blood flow and vascular density in murine models of PAD, including the db/db diabetic mouse model (26). Although no large-animal studies of treatment of CLI with NO-based therapeutics have been reported, a recent phase II clinical trial for safety and efficacy in which PAD patients were treated with oral sodium nitrite reported the drug to be well tolerated (32). High-dose sodium nitrite resulted in improved endothelial function, as seen by changes in flow-mediated vasodilation, along with improvements in overall physical functioning and pain, in diabetic patients (32).

In the current study we investigated a SR-nitrite formulation that was administered orally. Nitrite is metabolized fairly rapidly in vivo under conditions of tissue ischemia, and the SR-nitrite formulation utilized in the current study was selected to provide more continuous nitrite delivery to the ischemic limb. Safety was validated by stable circulating methemoglobin levels in the SR-nitrite-treated group. Pharmacokinetic analysis revealed a steady and significant rise in circulating nitrite, as well as nitrite delivery to the ischemic limb following treatment. Importantly, RSNO levels in the plasma and ischemic limb indicate that nitrite was converted to NO, as evidenced by increases in NO-bound protein. In the placebo-treated group, there was a significant reduction in NO and a trending decrease in nitrite levels in skeletal muscle of the ischemic compared the control limb. This reduction in NO
intermediates in the porcine CLI model is consistent with our finding that NO levels are significantly attenuated in skeletal muscle from CLI patients. The occlusion of large vessels presents a challenging scenario for a pharmacological agent, where the ischemic tissue requires extensive vessel formation to prevent necrosis. Preclinical studies using murine models of hindlimb ischemia have shown NO to be a proangiogenic factor (9, 28, 45). In the current study we evaluated whether nitrite therapy had an impact on collateralization and revascularization of larger vessels. Analysis of gastrocnemius skeletal muscle revealed a trend, but not statistically significant, increase in vessel density in the ischemic limb of the SR-nitrite-treated group. ABI measurements confirm that there was no improvement in ischemic limb blood pressure following SR-nitrite therapy. However, analysis of the angiogenic growth factor VEGF and markers for endothelial cell proliferation (CD31 and von Willebrand factor) indicated that SR-nitrite therapy did, in fact, promote signaling for vessel growth.

A primary limitation of this study is the extreme severity of CLI that is induced following implantation of a covered stent in combination with an AMPLATZER occluder in the external iliac artery. This procedure results in complete occlusion of the external iliac artery and initial absence of blood flow. The extreme reductions in hindlimb perfusion coupled with MetS result in an extremely severe model for testing therapies. Another potential limitation of our study is the delayed administration of SR-nitrite therapy at post-CLI day 14 and the relatively short duration of SR-nitrite therapy. Although therapy would have likely been more beneficial if administered at the time of CLI induction, we initiated SR-nitrite therapy in a clinically relevant timeline. Similarly, if we had administered the SR-nitrite beyond day 35, we may have observed improvements in hindlimb perfusion and ABI.

Uregulation of endothelial proliferation factors, such as VEGF, may not have resulted in arterial vessel formation and improved blood flow due to the severe model and delayed SR-nitrite therapy. It is not entirely surprising that sustained VEGF signaling observed in these studies (day 35) failed to result in endothelial proliferation. Recent studies suggest that prolonged elevations in VEGF-A can act as a negative regulator via a splicing mechanism to inhibit angiogenic signaling (39).

Finally, our data reveal that circulating and ischemic tissue levels of nitrite were significantly increased at 21 days following initial dosing, but we failed to observe statistically significant increases in plasma nitrite at the earlier time points. It is unclear exactly what circulating or tissue levels are required to exert cytoprotective and proangiogenic effects in animal models or humans. Therefore, it is conceivable that our dosing regimen did not result in therapeutic nitrite levels during the entire treatment period. Our dosing rationale was based on a phase II clinical trial to treat patients with PAD (32).

While the present study provides important new insights into nitrite therapy in CLI, additional studies are clearly required to more fully evaluate the therapeutic potential of SR-nitrite and other nitrite formulations in severe PAD and CLI to determine the potential efficacy of this NO-based therapeutic in patients with peripheral vascular disease.

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GRANTS

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DISCLOSURES

D. J. Lefer is on the scientific advisory board of Theravasc, Inc., which is currently developing sodium nitrite for the treatment of cardiovascular diseases. D. J. Lefer is a participant of a pending US patent filed on 14 October 2003 (patent no. 60/511,244) regarding the use of sodium nitrite in cardiovascular disease. D. J. Lefer is a participant of a pending US patent filed on 15 November 2007 (patent no. 61/003,150) regarding the use of nitrite salts in chronic ischemia.

AUTHOR CONTRIBUTIONS

D.J.P., J.M.B., L.P.B., C.C.C., I.L.P., T.T.G., and D.J.L. edited and revised the manuscript; D.J.P., J.M.B., T.T.G., and D.J.L. performed the experiments; D.J.P., J.M.B., K.N.I., and D.J.L. analyzed the data; D.J.P., J.M.B., I.L.P., and D.J.L. interpreted the results of the experiments; D.J.P., J.M.B., and D.J.L. figured the figures; D.J.P., J.M.B., and D.J.L. drafted the manuscript; D.J.P., J.M.B., L.P.B., J.W.C., C.C.C., I.L.P., T.T.G., and D.J.L. approved the final version of the manuscript.

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