Research Article

Vagus Nerve Stimulation Attenuates Acute Skeletal Muscle Injury Induced by Ischemia-Reperfusion in Rats

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Vagus nerve stimulation (VNS) has been shown to attenuate ischemia-reperfusion (I/R) injury in multiple organs. The present study aimed at investigating whether VNS could exert protective effects against I/R injury in the skeletal muscle. Male Sprague-Dawley rats were randomly divided into 3 groups: the control, I/R, and I/R+VNS groups. The skeletal muscle I/R (SMI/R) model was induced by occlusion of the left femoral artery for 2.5 hours followed by reperfusion for 2 hours. The vagal nerve trunk was separated, and VNS was performed during the whole I/R process. The intensity of VNS was optimized in each rat to obtain a 10% reduction in the heart rate relative to the value before stimulation. After the experiment, the blood sample and left gastrocnemius muscle tissues were collected for histological examination, biochemical analysis, and molecular biological detection. During the I/R process, VNS significantly reduced cellular apoptosis, necrosis, and inflammatory cell infiltration compared to sham VNS. The VNS treatment also decreased the inflammatory response, alleviated oxidative stress, and improved vascular endothelial function (p < 0.05 for each). In contrast, the I/R group showed an opposite effect compared to the control group. The present study indicated that VNS could protect against SMI/R injury by suppressing excessive inflammation, alleviating oxidative stress, and preserving vascular endothelial function.

1. Introduction

Skeletal muscle ischemia-reperfusion (SMI/R) injury is a common disease in clinical practice. It often influences the function of the skeletal muscle and can even be life-threatening [1]. However, there are few effective treatments for SMI/R [2]. Thus, novel effective therapies are needed to prevent SMI/R injury. Numerous studies have demonstrated that the pathogenesis of SMI/R injury is associated with inflammation responses and oxidative stress [3]. Vagus nerve stimulation (VNS) has been shown to exert anti-inflammatory and antioxidative effects [4–6]. Most recently, VNS has been demonstrated to improve I/R injury in multiple organs, including the heart, brain, and kidney [7–9]. VNS applied during the I/R process could reduce the infarct area and improve the prognosis. More importantly, our clinical study has shown that transcutaneous VNS can markedly attenuate myocardial I/R injury in acute myocardial infarction patients undergoing primary percutaneous coronary intervention [10]. However, the protective effects of VNS against SMI/R injury remain unknown. In the present study, using an acute SMI/R model in rats, we aimed to demonstrate the protective effects of VNS against SMI/R injury and further explore the potential mechanisms.

2. Materials and Methods

2.1. Animals and Experimental Groups. Healthy male Sprague-Dawley rats (250-300 g) were included in this study,
and all animals were supplied by the Experimental Animal Center of Renmin Hospital of Wuhan University. The study conformed to the Guidelines for the Care and Use of Laboratory Animals published by the US National Institutes of Health (NIH Publication, revised 1996). All animal studies were reviewed and approved by the Renmin Hospital of Wuhan University Animal Care and Use Committee (ethics clearance number was WDRM. 20180308).

Rats were randomized into 3 groups and received the following treatments: sham operation (control group, \( n = 6 \)), SMI/R with sham VNS (I/R group, \( n = 6 \)), and SMI/R with VNS (I/R+VNS group, \( n = 6 \)). Detailed study protocol is summarized in Figure 1(a).

### 2.2. Acute SMI/R Model

Rats were anesthetized with 2% pentobarbital sodium (40 mg/kg body weight) by intraperitoneal injection. Surface electrocardiography was performed with a PowerLab data acquisition system (8/35, ADInstruments, Bella Vista, Australia). The left femoral artery was exposed by blunt dissection and a pair of atrumatic microvascular clamps were placed (in the control group, only femoral artery exposure was performed). The presence of pulsation under the clamp was checked. After 2.5 h of ischemia, the microvascular clamps were removed and the left hind limb received 2 h of reperfusion as previously described [11].

### 2.3. VNS

The left cervical vagal trunk was isolated as a stimulating target (see Figure 1(b)). Continuous stimulation (20 Hz, 0.1 ms in duration, square waves) was delivered by a stimulator (S20, Jinjiang, Chengdu City, China) through a pair of Teflon-coated silver hooks (0.1 mm in diameter) on the cervical vagal trunk. The stimulation level was defined as the voltage level sufficient to slow the sinus rate or atrioventricular (AV) conduction at 10%, as previously described [12]. The VNS threshold was determined once again prior to each hour of stimulation.

### 2.4. Blood and Tissue Sample Collection

After the entire experimental progress, blood samples were collected from the inferior vena cava. Serum was collected by centrifugation at 3,000 rpm, for 15 min. Tissue specimens were collected from the first half of the left gastrocnemius muscle venter. Each tissue specimen was separated into three parts. The major part (about \( 5 \times 5 \times 10 \text{ mm}^3 \)) was used for histological analysis. The two minor parts (about \( 2 \times 2 \times 2 \text{ mm}^3 \)) were used for biochemical and molecular biological analysis. All blood and tissue samples for biochemical and molecular biological analysis were stored at -80°C until use.

### 2.5. Histological Examination

The skeletal muscle tissue samples were submerged in 10% paraformaldehyde solution for 48 h, dehydrated sequentially in an ascending gradient of ethanol, and rinsed in xylene. Then, the tissues were embedded in paraffin. Sections of 5 μm thick were stained with hematoxylin and eosin (H&E) and then examined under a light microscope.

The muscle injury area was estimated in each gastrocnemius muscle section under a microscope according to Yin et al. [13].

Terminal deoxynucleotidyl transferase dUTP nick end labeling (TUNEL) staining was performed with a commercially available kit (Roche Biochemicals, Mannheim, Germany) according to the manufacturer’s protocol. Tissue sections were deparaffinized and then hydrated with ethanol. After hydration, the sections were treated with protease K, rinsed with phosphate-buffered saline (PBS), and then incubated with TUNEL reaction reagents. After washing with PBS, the sections were treated with 4′,6-diamidino-2-phenylindole (DAPI) and incubated in a dark environment at room temperature. Stained sections were analyzed using a fluorescence microscope (Nikon DS-U3, Japan). Cells with nuclei containing irregular green particles were defined as TUNEL-positive cells. Cell death was viewed and detected in three random fields of each muscle section and averaged. The TUNEL-positive cell ratio was recorded.

### 2.6. Analysis of Serum Creatine Kinase (CK) and Lactate Dehydrogenase (LDH) in Serum Levels

Serum CK and LDH levels, which reflect skeletal muscle injury severity, were assessed using commercial kits (Changchun Huili Biotech Co. Ltd, China) according to the manufacturer’s protocol.

### 2.7. Measurement of Malondialdehyde (MDA), Myeloperoxidase (MPO), and Superoxide Dismutase (SOD) Content Levels in Serum and Tissues

Tissue and serum samples from the control, I/R, and I/R+VNS groups were homogenized. The tissue MPO activity level, the tissue and serum MDA concentration, and SOD activity levels were assayed using commercially available kits (Nanjing Jiancheng Bioengineering Institute, Nanjing City, China) according to the manufacturer’s protocol. The samples after reaction were detected by a spectrophotometer. The maximum absorbance values determined for MDA, MPO, and SOD were 532 nm, 460 nm, and 550 nm, respectively.

### 2.8. Western Blot Analysis

Frozen skeletal muscle tissues were lysed with a RIPA lysis buffer (Aspen Biotechnology, Wuhan, China), supplemented with phenylmethanesulfonyl fluoride (Aspen). After homogenization, the supernatant was collected for protein concentration determination using a BCA protein assay kit (Aspen). Equal amounts of protein solution from the homogenates were subjected to SDS-PAGE and then transferred onto a nitrocellulose membrane. After the membrane was blocked with 5% fat-free milk, it was incubated with rabbit anti-endothelial nitric oxide synthase antibody (eNOS, 1:1000 dilution,Abcam, Cambridge, UK), rabbit anti-intercellular adhesion molecule 1-antibody (ICAM-1, 1:500 dilution, Biorbyt, UK), rabbit anti-intercellular adhesion molecule-1 antibody (VCAM-1, 1:1000 dilution, Abcam, Cambridge, UK), rabbit anti-interleukin 1 beta antibody (IL-1β, 1:1000 dilution, Abcam, Cambridge, UK), rabbit anti-IL-6 antibody (1:1000 dilution, Affinity, San Francisco, USA), and rabbit anti-tumor necrosis factor alpha antibody (TNF-α, 1:1000 dilution, Proteintech Group, Inc. Wuhan, China). After washing three times with Tris-buffered saline containing 1% Tween (TBST), the membranes were incubated with HRP-goat anti-rabbit antibody (Aspen) at room temperature for 30 min and then washed four times with TBST. The protein bands were visualized by the...
2.9. qRT-PCR. Total RNA was extracted from the frozen skeletal muscle tissues using a Trizol reagent (Invitrogen Life Technologies) according to the manufacturer’s protocol. First strand cDNA was synthesized using PrimeScript™ RT Reagent Kit with gDNA Eraser (TaKaRa Bio Inc.). Candidate gene expression levels were measured using an RT-PCR thermocycler (StepOne™ Life Technologies) with the following specific primers: IL-1β, forward: 5′-GTGG CAGCTACCTATGTCTTGCG-3′, reverse: 5′-CCACCTTGGTGCGCTATGTTCTGT-3′; IL-6, forward: 5′-TGGAGTTCC GTTTCACCTTG-3′, reverse: 5′-GGTCCTTAGCCACT CTTTGTGTCTC-3′; and TNF-α, forward: 5′-CACCACGCTCT TCTGTCTACTG-3′, reverse: 5′-GCTACGGGCCCTGTAC TCG-3′. The mRNA level for each target gene was calculated using the Delta-Delta-CT method and normalized to the β-actin mRNA level from the same sample. The primers for β-actin were as follows: forward: 5′-CGTGTGACATCC GTAAAGACCTC-3′, reverse: 5′-TAGGAGCCAGGCA GTAATCT-3′.

2.10. Statistical Analysis. Continuous variables are expressed as the mean ± SD and were analyzed by one-way ANOVA. All data were analyzed using GraphPad Prism version 7.0 software (GraphPad Software, Inc. San Diego, CA), and two-tailed p ≤ 0.05 was considered significant.

3. Results

3.1. VNS Significantly Alleviated SMI/R Injury. As shown in Figure 2(a), skeletal muscle tissues in the I/R group showed muscle degeneration, necrosis, sarcoplasmic dissolution, and neutrophil infiltration. The extent of these changes was alleviated in the I/R+VNS group. Figure 2(b) shows that the gastrocnemius muscle injury area in the high-power field (HPF) was highest in the I/R group, significantly lower in the I/R+VNS group, and lowest in the control group.

Figure 2(c) shows that the CK and LDH levels were significantly higher in the I/R group than in the control group (CK, 5060.31 ± 847.02 vs. 233.13 ± 98.01 U/L, p < 0.05; LDH, 847.96 ± 120.20 vs. 298.50 ± 73.10 U/L, p < 0.05). The CK and LDH levels were significantly attenuated by the VNS treatment (CK, 1849.44 ± 456.94 vs. 5060.31 ± 847.02 U/L, p < 0.05; LDH, 625.65 ± 81.51 vs. 847.96 ± 120.20 U/L, p < 0.05).

TUNEL staining was used to detect skeletal muscle cell apoptosis among the three groups. The percentage of TUNEL-positive cells was significantly increased in the I/R group compared with the control group (32.83 ± 4.62% vs. 1.67 ± 1.21%, p < 0.05). In the I/R+VNS group, the percentage of TUNEL-positive cells was significantly decreased compared with that in the I/R group (13.83 ± 2.59% vs. 32.83 ± 4.62%, p < 0.05) (see Figure 3).

3.2. VNS Significantly Mitigated the Inflammatory Response in the Skeletal Muscle after I/R. As shown in Figure 4, the mRNA and protein expression levels of proinflammatory factors IL-1β, IL-6, and TNF-α were markedly increased in the I/R group compared with the control group (mRNA expression: IL-1β, 2.92 ± 0.44 vs. 1.00 ± 0.08; IL-6, 2.80 ± 0.20 vs. 1.00 ± 0.13; and TNF-α, 3.88 ± 0.31 vs. 1.00 ± 0.17, p < 0.05 for each. Protein expression: IL-1β, 4.03 ± 0.58 vs. 1.00 ± 0.42; IL-6, 3.20 ± 0.32 vs. 1.00 ± 0.31; and TNF-α, 4.40 ± 0.62 vs. 1.00 ± 0.6, p < 0.05 for each). VNS significantly mitigated the increased expression of levels of inflammatory markers (mRNA expression: IL-1β, 1.71 ± 0.11 vs. 2.92 ± 0.44; IL-6, 1.76 ± 0.09 vs. 2.80 ± 0.20; and TNF-α, 2.42 ± 0.15 vs. 3.88 ± 0.31, p < 0.05 for each. Protein expression: IL-1β, 2.29 ± 0.51 vs. 4.03 ± 0.58; IL-6,
1.83 ± 0.43 vs. 3.20 ± 0.32; and TNF-α, 2.70 ± 0.74 vs. 4.40 ± 0.62, p < 0.05 for each).

3.3. VNS Significantly Attenuated Oxidative Stress in Skeletal Muscle after I/R. The MPO activity in skeletal muscle tissues was used to indicate the neutrophilic infiltration severity. I/R injury significantly increased the MPO activity compared with the sham operation (0.40 ± 0.06 vs. 0.13 ± 0.02 U/g, p < 0.05). In contrast, VNS significantly alleviated the increase in MPO activity after I/R (0.25 ± 0.04 vs. 0.40 ± 0.06 U/g, p < 0.05) (see Figure 5(c)). The changes in the MDA and SOD concentrations in skeletal muscle tissues are shown in Figures 5(a)-(b). As a biomarker of oxidative stress, the MDA level was significantly increased in tissues after I/R (1.29 ± 0.14 vs. 0.54 ± 0.08 nmol/mg protein, p < 0.05). In contrast, as a biomarker of antioxidant activity, the SOD was...
significantly decreased in tissues after I/R (39.26 ± 7.85 vs. 118.85 ± 23.00 U/mg protein, p < 0.05). However, VNS during I/R markedly mitigated these changes (MDA, 0.92 ± 0.07 vs. 1.29 ± 0.14 nmol/mg protein; SOD, 68.31 ± 14.46 vs. 39.26 ± 7.85 U/mg protein, p < 0.05). Similar variation trends were observed for the serum MDA and SOD levels (see Figures 5(d)-5(e)). Compared to the control group, the I/R group showed significantly different serum MDA and SOD levels (MDA, 8.58 ± 0.88 vs. 4.32 ± 0.62 nmol/mL; SOD, 175.14 ± 13.10 vs. 383.93 ± 26.28 U/mL, p < 0.05 for both), while VNS markedly attenuated these changes (MDA, 6.19 ± 0.80 vs. 8.58 ± 0.88 nmol/mL; SOD, 305.37 ± 14.76 vs. 175.14 ± 13.10 U/mL, p < 0.05 for both).

3.4. VNS Significantly Protected Vascular Endothelial Function in the Skeletal Muscle after I/R. Endothelial function was evaluated by the expression of eNOS, ICAM-1, and VCAM-1. Figure 6(b) shows that I/R significantly decreased the expression level of eNOS and increased the expression levels of ICAM-1 and VCAM-1, while VNS exerted a protective effect on endothelial function and relieved the above changes. The relative expression of eNOS, ICAM-1, and VCAM-1 in the control and I/R groups were 1.00 ± 0.09 vs. 0.25 ± 0.10, 1.00 ± 0.28 vs. 5.15 ± 0.77, and 1.00 ± 0.20 vs. 3.82 ± 0.59, respectively, with p < 0.05 for all. VNS significantly relieved the changes described above (0.59 ± 0.16 vs. 0.25 ± 0.10, 3.40 ± 0.83 vs. 5.15 ± 0.77, and 2.14 ± 0.61 vs. 3.82 ± 0.59 for eNOS, ICAM-1, and VCAM-1, respectively, p < 0.05 for all).

4. Discussion

In the present study, we provide novel evidence that VNS during SMI/R injury could ameliorate skeletal muscle injury, as shown by alleviated cellular apoptosis, degeneration and inflammatory cell infiltration, and reduced serum CK and LDH levels compared to sham VNS. The underlying mechanisms of this protective effect involve inhibiting excessive inflammation and oxidative stress and protecting endothelial function. To the best of our knowledge, this is the first study to apply VNS to treat SMI/R injury.

It is well known that the innervation of the autonomic nervous system plays a vital role during organic I/R injury, including SMI/R injury. Sympathetic nerves are distributed in the adventitia of arteries in the skeletal muscle [14]. Increasing the sympathetic tone will lead to enhanced noradrenaline release, which results in vasoconstriction via direct activation of the α-receptors and inhibition of
vasodilating neuropeptides. Ischemia and hypoxia are common sympathoexcitatory stresses. This sympathetic vasoconstrictive effect might further exaggerate SMI/R injury [15]. Povlsen and Sirsjo have reported that treatment with guanethidine, a sympathetic nerve blocker, during reperfusion in SMI/R leads to a better prognosis [16]. Increased vagal tone might offset the sympathetic vasoconstrictive effect and protect organs from I/R injury. There are also emerging studies of this potential therapy for I/R injury. In a rat model, Jiang et al. reported that VNS treatment during cerebral I/R significantly reduced I/R injury [17]. The heart muscle is histologically similar to the skeletal muscle, and in a cardiac I/R model, VNS significantly reduced reperfusion arrhythmias and infarct size [8]. Our previous study also showed an analogous protective effect of VNS against myocardial I/R injury [18]. Such effects suggest that VNS might be a potential approach for treating SMI/R. The results of the present study are consistent with those of previous studies and demonstrate that VNS can attenuate SMI/R injury.

Currently, it is recognized that the pathophysiological mechanisms of I/R injury in the skeletal muscle include inflammation, oxidative stress, vascular endothelial damage, calcium overload, and mitochondria damage [19–22]. Corrick et al. reported that the administration of dexamethasone, a classic anti-inflammatory drug, at the onset of reperfusion ameliorated the structural and functional damage in the skeletal muscle [23], indicating the important role of inflammation in SMI/R injury. Acetylcholine (Ach) is an anti-inflammatory substance through cholinergic anti-inflammatory pathways [24]. VNS has been demonstrated to stimulate the release of Ach and might be a potential anti-inflammatory treatment in different diseases. Jonge et al. have reported that VNS could attenuate macrophage activation and suppress inflammation [25]. Similar results have been reported by Koopman et al., indicating that VNS could inhibit proinflammatory cytokine production and attenuate inflammatory disease [26]. Several researchers have applied the vagal anti-inflammatory effect in I/R injury and achieved excellent effects. Inoue et al. reported that VNS significantly reduced the expression of proinflammatory cytokines in a renal I/R rat model [7]. Similarly, our data indicate that VNS significantly reduced the inflammatory cytokine levels compared with sham VNS.

Reperfusion induced oxidative stress can promote skeletal muscle cell apoptosis [27]. It is known that inhibiting oxidative stress can effectively alleviate I/R injury. One study has shown that VNS can reduce oxidative stress in a
cerebral I/R rat model [17]. Our previous study indicated that VNS markedly reduced reactive oxygen species (ROS) production in a myocardial I/R canine model [18]. These data show that VNS could prevent I/R injury via an antioxidative stress effect. MPO is involved in the generation of ROS. The balance between prooxidant biomarkers (MDA) and antioxidant biomarkers (SOD) represents the activity of oxidant stress [28]. In the present study, our data suggest that VNS significantly decreased tissue MPO activity, reduced the concentration of MDA, and increased SOD activity in both tissue and serum. Therefore, we suggest that antioxidative activity may be one of potential mechanisms underlying the protective effect of VNS against SMI/R injury.

Vascular endothelial dysfunction is another mechanism involved in SMI/R injury [29]. Endothelial activation, which is defined by increased expression of cell surface adhesion molecules such as ICAM-1 and VCAM-1, was the main manifestation of vascular endothelial dysfunction [30]. The inflammatory response could activate the vascular endothelium, promote the production of proinflammatory cytokines (such as IL-6 and TNF-α), and increase the expression of ICAM-1 and VCAM-1. A previous study has shown that cholinergic agonists could suppress endothelial cell activation, as confirmed by decreased ICAM-1 expression [31]. Our results show that VNS relieved the increased expression of ICAM-1 and VCAM-1 induced by SMI/R injury. Nitric oxide (NO) is a bioactive molecule that helps dilate the blood vessels. It has been proven that NO could inhibit inflammatory cell adhesion and limit endothelial activation [32]. The expression of eNOS is essential for the production of NO and integrity of the vascular endothelium [33]. Yoshiizumi et al. reported inhibiting eNOS expression could lead to endothelial dysfunction in human umbilical vein endothelial cells [34]. Previous studies have shown that increased eNOS expression leads to improvement of the endothelial function, which could improve the prognosis of SMI/R injury [11]. Li et al. reported that chronic VNS increased the expression of eNOS in ovariectomized rats [35]. In the present study, VNS significantly increased the expression of eNOS compared with sham VNS. Together with the downregulated expression of ICAM-1 and VCAM-1 and the increased eNOS expression results, our data indicated that VNS could protect the skeletal muscle from I/R injury by preserving endothelial function.

SMI/R injury is a common clinical condition. Until now, effective drug interventions to address this pathological state have been limited. The present study provides evidence that VNS could markedly reduce skeletal muscle tissue injury and cell apoptosis induced by I/R. Recently, stimulating the auricular branch of the vagus nerve has been proven to be safe and effective for achieving a similar effect as cervical VNS, which might overcome the shortcomings of conventional VNS [10, 36]. With further research to validate its safety and practicability, noninvasive VNS might become a novel technology to treat SMI/R injury in patients (see Figure 7).

4.1. Study Limitations. First, different stimulation sites and parameters have been shown to exert distinct therapeutic effects. In this study, we only verified the current stimulation parameters in left-side VNS. Further studies will aim

Figure 6: VNS protects vascular endothelial function. (a) Representative examples of eNOS, ICAM-1, and VCAM-1 expression in different groups. (b) Relative expression levels are shown as the ratio between the target proteins and β-actin expression levels. *p < 0.05 vs. control group; #p < 0.05 vs. I/R group. eNOS: endothelial nitric oxide synthase; ICAM-1: intercellular cell adhesion molecule-1; VCAM-1: vascular cell adhesion molecule-1.
to contrast bilateral VNS and explore the best stimulation parameters. Second, although we revealed several potential mechanisms by which VNS protects against SMI/R injury, the exact mechanism remains to be explored. Third, we only investigated the acute impact of VNS on SMI/R injury. Long-term effects should be verified in future studies. Fourth, we only measured the expression of eNOS as a parameter to reflect the endothelial function. The eNOS activity and NO production can be better to reflect the endothelial function. The effect of VNS on these parameters will be investigated in our further studies.

5. Conclusions

In conclusion, our data suggest that VNS could play a protective role in I/R-induced skeletal muscle injury. Its potential mechanisms may involve suppressing excessive inflammation, alleviating oxidative stress, and preserving vascular endothelial function. Although further studies are needed to validate its safety and practicability, VNS might provide a novel treatment for patients with SMI/R injury.

Data Availability

The data used to support the findings of this study are available from the corresponding author upon request.

Conflicts of Interest

The authors declare that there is no conflict of interest regarding the publication of this paper.
Authors’ Contributions
Dr. Zhang and Dr. Li contributed equally to this study.

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