Impact of intracoronary nicorandil before stent deployment in patients with acute coronary syndrome undergoing percutaneous coronary intervention

XINGLI XU¹, XIAOLING LIU¹, LIWEN YU¹, JING MA¹, SUFANG YU² and MEI NI¹

¹Key Laboratory of Cardiovascular Remodeling and Function Research, Chinese Ministry of Education, Chinese National Health Commission and Chinese Academy of Medical Sciences, The State and Shandong Province Joint Key Laboratory of Translational Cardiovascular Medicine, Department of Cardiology, Qilu Hospital of Shandong University, Jinan, Shandong 250012;
²Department of Neurology, The Fourth People’s Hospital, Liaocheng, Shandong 252002, P.R. China

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Abstract. The present study aimed to clarify the effect of bolus intracoronary nicorandil on inflammatory, oxidative and adherent indicators in patients with acute coronary syndrome (ACS) undergoing percutaneous coronary intervention (PCI). This randomized controlled trial (RCT) was performed to detect the inflammation and oxidative stress in intracoronary blood both before and after PCI. In total, 65 consecutive patients undergoing PCI were classified into a nicorandil therapy group (n=32) or a placebo group (n=33). All procedures were performed at Shandong University Qilu Hospital, China, during the period from March, 2016 to May, 2017. Intracoronary blood from patients who received nicorandil therapy during PCI showed no change in soluble CD40 ligand (sCD40L) concentration (1.86±0.08 vs. 1.90±0.09 ng/ml, P=0.12) but a significant increase was noted in the control group (1.87±0.17 vs. 2.82±0.26 ng/ml, P<0.01). This indicated a relative reduction in sCD40L level after PCI in the nicorandil group. We further demonstrated an increase in superoxide dismutase (SOD) activity (29.37±0.81 vs. 31.03±0.60 U/ml, P=0.001) and a reduction in lipid peroxidation (3.84±0.99 vs. 4.23±0.13 U/ml, P=0.001) in the nicorandil group but observed no change in the placebo group. ICAM-1 levels showed no change in the nicorandil group (69.54±6.89 vs. 72.01±8.25 ng/ml, P=0.83) but a significant increase in the control group after PCI in intracoronary blood (56.57±4.96 vs. 76.81±6.88 ng/ml, P=0.002). No changes were found in hs-CRP, TNFα and sVCAM-1 levels in coronary blood for both groups before and after PCI in ACS patients. Our findings demonstrate that intracoronary bolus nicorandil therapy has a significant effect on the inhibition of inflammatory indicators and oxidative stress in patients with ACS during PCI. This suggests a possible medical application of nicorandil for reducing inflammation and oxidative stress.

Introduction

Percutaneous coronary interventions (PCIs) are widely performed as an invasive intervention to treat patients with acute coronary syndromes (ACSs) including non-ST-elevation ACS [including unstable angina (UA)] and non-ST-elevation myocardial infarction (NSTEMI) and ST-elevated myocardial infarction (STEMI). It is well-established that PCI induces elevated levels of inflammatory mediators and pathophysiological levels of reactive oxygen species (ROS) production, which leads to an occurrence of restenosis, stent thrombosis and other adverse cardiovascular events in patients with ACS after PCI (1-4).

Inflammatory markers have been implicated as the most important risk indicators in the development of restenosis and may be predictive of poor cardiovascular outcomes in ACS.
patients undergoing coronary angiography (4-7). The concentration of inflammatory cytokines also correlates with the occurrence of stent restenosis. Furthermore, oxidative stress is involved in the pathogenesis of cardiovascular diseases, which is characterized by an imbalance between the generation of ROS and the capacity of the intrinsic antioxidant defense system (8). Research has documented that oxidative stress indicators released from stenotic and symptomatic lesions in patients undergoing PCI participate in the development of new lesions, and increase the risk of major adverse cardiovascular events (MACE) such as cardiovascular death, myocardial infarction (MI) and stroke (9–13).

Nicorandil is a hybrid agent with distinctive pharmacological function as an ATP-sensitive potassium channel (K_{ATP}) opener and a nitric oxide donor. The mechanism of nicorandil includes not only dilation of both the macrovascular and microvascular systems but also has potential cardio-protective effects. Intracoronary (IC) injection of nicorandil has been reported to be safe for patients with coronary artery disease (CAD) during PCI (14–18). Additionally, it was found to improve the prognosis of patients with acute heart failure (AHF) (16), and to ameliorate both early functional deflection (17) and long-term complications of acute myocardial infarction (AMI). However, the effects of intracoronary nicorandil injection on inflammation and oxidative stress remain unclear in patients with ACS undergoing PCI as no data are available at present. Thus, the present study was performed to evaluate the feasibility and efficacy of IC nicorandil injection for the improvement of inflammation and oxidative stress in patients with ACS undergoing PCI.

Materials and methods

Study population. This single-blinded, randomized prospective and clinical trial was performed at the Department of Cardiology, Qilu Hospital, Shandong University, Jinan, Shandong, China. Patients were eligible if they were age 30 years or older and admitted to our institution for ACS. The diagnosis of ACS was based on clinical symptoms, electrocardiographic changes compatible with AMI, and elevation of cardiac biomarkers (18). This study enrolled patients with AMI and unstable angina pectoris. Clinical symptoms of ACS were as follows: Angina pectoris including chest pain with or without radiating to the neck, jaw, upper abdomen or shoulder. AMI was defined as episodes of chest pain persisting >30 min and <24 h, ST-segment elevation in at least two continuous electrocardiogram leads, and more than two-fold creatine kinase (CK) elevation above the maximum peak in the normal range (19). The diagnosis of unstable angina pectoris was based on typical precordial chest pain at rest, angiographic evidence of a stenosis ≥75% according to the American Heart Association (AHA) classification (11) and no elevation of cardiac biomarkers.

Exclusion criteria were as follows: Age >90 years; patients with history of coronary artery bypass grafting (CABG); systolic blood pressure (BP), 80 mmHg; known allergy to aspirin, clopidogrel, ticagrelor or nicorandil; bleeding history within the prior 3 months; cancer or other severe comorbidity affecting life expectancy. NSTEMI patients are not eligible for stent implantation, thus they were also excluded. Finally, 65 consecutive participants with a diagnosis of ACS were recruited to the trial after providing written informed consent for angiography, PCI and blood extraction. All patients were admitted to our hospital between March 2016 and May 2017. This study was approved by the Qilu Hospital Ethics Committee (KYLL-2015266). All procedures were performed as standard interventional techniques following current guidelines at the time of intervention.

Procedures. Sixty-five patients with ACS were randomly divided into two groups: A nicorandil treatment group and a control group. The 32 patients in the nicorandil group received an IC bolus of nicorandil (2 mg/2 ml) before reperfusion with PCI and the 32 patients in the control group received the same volume of physiological saline. PCIs were performed due to stenosis-related coronary artery. Next, 3,000 units of heparin was injected after the achievement of arterial access, and adjunctive injections of 7,000 units were needed for patients undergoing PCI. IC administration of nitroglycerin was used to achieve the maximal vasodilation for all patients before the initial and final angiograms. PCI was performed following standard procedures.

A total dose of 12 mg nicorandil was dissolved in 12 ml of 0.9% saline. Two milliliters of nicorandil in the nicorandil group and 2 ml saline in the control group were administered, respectively, by IC bolus injection before percutaneous transluminal coronary angioplasty (PTCA). All other procedures were the same for both treatment groups. The cardiologist in-charge performed PCI and IC injection of nicorandil or saline in the catheter laboratory at the appropriate time. No angiographical residual stenosis was detected in any patients and there were no deaths during PCI. No follow-up data, such as complications, mortality and long-term effect of nicorandil, were available in this study. After all procedures, patients were maintained on aspirin (100 mg once daily) plus clopidogrel (75 mg once daily) or ticagrelor (90 mg twice daily) for at least 1 year after metal stent placement followed by the physician’s specific directions.

Collection and preservation of coronary blood. Approximately 10 ml of intracoronary blood was collected from the distal coronary stenosis with a Finecross micro-catheter (Asahi Intecc Co., Ltd.) before the injection of nicorandil or saline prior to PTCA, and immediately after the stent placement in patients undergoing PCI. The blood samples were collected in vacutainer anticoagulant tubes at room temperature for ~30 min and then centrifuged for 10 min at 1,000 x g. Plasma and serum were aliquoted and stored at -80°C until further analysis.

Measurement of SOD and MDA. Superoxide dismutase (SOD) was examined using an SOD determination kit (cat. no. 19160; Sigma-Aldrich; Merck KGaA) following the manufacturer’s instructions. Each sample was measured with three replicates of four samples: Sample, Blank 1, Blank 2 and Blank 3. The Sample subset consisted of 20 µl sample solution, 200 µl of WST working solution and 20 µl of enzyme working solution. Then the sample solution was substituted by 20 µl of enzyme working solution as Blank 1. Blank 2 consisted of 20 µl sample solution, 200 µl of WST working solution and 20 µl of dilution solution.
Results

Patient baseline characteristics. In this study, a total of 65 consecutive patients were enrolled with 32 randomly allocated to the nicorandil and 33 to placebo groups. The baseline characteristics of the two groups are shown in Table I. There were no significant differences in age, sex, BMI, incidence of coronary risk factors, serum creatinine or urea nitrogen levels, ejection fraction (EF), previous MI history, culprit lesions or quantitative angiographic data between the two groups. Clinically, 5 patients (15.6%) and 3 patients (9.1%) were diagnosed with myocardial infarction (MI) in the nicorandil group and placebo group, respectively. Other patients were diagnosed with unstable angina (UA). There were no significant differences in the use of oral medications at discharge among all patients.

Outline of the PCI procedures. Coronary angiography was performed to visualize the severity of coronary stenosis. The subsequent treatment of PTCA was followed up if the residual diameter stenosis was greater than or equal to 75% of the coronary artery stenosis diameter. The Finecross micro-catheter reached the distal coronary stenosis guided by a coronary guide-wire, and 10 ml of coronary blood was collected before PCI. Then 2 ml nicorandil (1 mg/ml) or placebo (2 ml of 0.9% saline) was IC bolus injected into coronary ostia. After nicorandil (or placebo) injection, PCI was performed according to standard techniques. Another 10 ml of coronary blood was collected immediately after stent implantation (Figs. 1 and S1).

Effect of nicorandil on markers of inflammation. There was a significant increase in soluble CD40 ligand (sCD40L) levels in the plasma of post-PCI compared with pre-PCI in the control group (1.87±0.17 vs. 2.82±0.26 ng/ml, P<0.001), but no difference in the nicorandil group in both pre- and post-PCI (1.86±0.08 vs. 1.90±0.09 ng/ml, P=0.12) (Fig. 2A). Compared with post-PCI in the control group, there was a significant reduction in sCD40L levels in the nicorandil group (P=0.001) (data not shown). The amount of change (Δ value) for sCD40L was significantly increased in the control group with no difference in the nicorandil group (0.04±0.04 vs. 0.95±0.23 ng/ml, P=0.001; Fig. 2A). No significant change was found in pre- and post-PCI levels of TNFα in the nicorandil group (5.90±1.84 vs. 5.3±2.73 pg/ml, P=0.43) and control group (6.13±0.58 vs. 7.08±0.95 pg/ml, P=0.29) (Fig. 2B). In comparison, there was no difference between pre-PCI and post-PCI in levels of IC plasma hs-CRP in the nicorandil group (5.16±1.05 vs. 6.13±1.13 mg/ml, P=0.054) and in the control group (5.98±1.01 vs. 6.64±1.12 mg/ml, P=0.458) (Fig. 2C). The Δ value of both TNFα (-0.60±0.75 vs. 0.95±0.84 pg/ml, P=0.405) and hs-CRP (0.98±0.49 vs. 0.66±0.88 mg/ml, P=0.246) remained statistically unchanged (Fig. 2B and C). Thus, the inflammatory sCD40L marker was significantly improved during PCI due to the intracoronary bolus injection of nicorandil.

Effect of nicorandil on indicators of oxidative stress. SOD and MDA indicators were measured to investigate the effect of nicorandil on oxidative stress. SOD was significantly increased in the plasma of post-PCI compared with pre-PCI in nicorandil group (29.37±0.81 vs. 31.03±0.60 U/ml, P<0.001), and no difference was observed between time-points in the control group (29.29±0.68 vs. 29.58±0.73 U/ml, P=0.519) (Fig. 3A). Additionally, MDA was significantly increased in
plasma of post-PCI compared to pre-PCI in the nicorandil group (3.84±0.10 vs. 4.230±0.13 mmol/ml, P=0.001), and no difference was observed between time-points in the control group (3.78±0.11 vs. 3.85±0.13 mmol/ml, P=0.402) (Fig. 3B). Surprisingly, the ratio of SOD and MDA was unchanged in four time-point subgroups of both the nicorandil (7.84±0.33 vs. 7.58±0.30, P=0.241) and control (7.92±0.27 vs. 7.95±0.35, P=0.888) group (Fig. 3C). There also was no difference in the Δ value of SOD (1.28±0.39 vs. 0.29±0.45 U/ml, P=0.328) in the control vs. nicorandil group, but a significant increase in levels of MDA (-0.70±0.26 vs. 0.07±0.08 mmol/ml, P<0.001) in the control group compared with the nicorandil group (Fig. 3A and B).

These results indicated that nicorandil may relieve oxidative stress via increased levels of SOD, but the levels of MDA were also subsequently increased. The possible change in ratio between SOD and MDA during PCI may be based on operational injury and the delayed reaction of the plasma levels to oxidative stress indicators.

Effect of nicorandil on angiokinetic factors. To further explore the effects of bolus injection of nicorandil during PCI, we also measured two different functional angiokinetic factors, ICAM-1 and sVCAM-1 post-PCI compared with pre-PCI. The results showed no significant differences in ICAM-1 in the nicorandil group (64.79±3.63 vs. 63.80±5.85 ng/ml, P=0.827).
Figure 1. Outline of the study procedures. Patients with UA or STEMI underwent PCI performed according to the standard techniques and using 1 of 2 strategies: i) PCI with nicorandil IC bolus injection before PTCA or ii) PCI followed by the same volume of 0.9% saline IC bolus injection before PTCA. PCI, percutaneous coronary intervention; PTCA, percutaneous transluminal coronary angioplasty; UA, unstable angina; STEMI, ST-elevated myocardial infarction.

Figure 2. Dynamics of the inflammatory response pre-PCI and post-PCI, and the difference in values between the two subgroups in the nicorandil and control groups. (A) Levels of sCD40L before and after PCI in the groups and subgroups, and Δ values of sCD40L of post-PCI compared with pre-PCI in the two groups. (B) Levels of TNFα before and after PCI in the groups and subgroups, and Δ values of TNFα of post-PCI compared with pre-PCI in the two groups. (C) Levels of hs-CRP before and after PCI in the groups and subgroups, and different values of hs-CRP of post-PCI compared with pre-PCI in two groups. Values are expressed as mean ± SD; Δ values are expressed as mean ± SD; **P<0.01, ***P<0.001. PCI, percutaneous coronary intervention; sCD40L, soluble CD40 ligand; TNFα, tumor necrosis factor α; hs-CRP, high-sensitivity C-reactive protein.
However, levels of ICAM-1 in the control group were increased after PCI (66.30±4.17 vs. 75.24±5.46 ng/ml, P=0.002) (Fig. 4A). There was also no change in levels of sVCAM-1 in the nicorandil group (4.85±0.61 vs. 5.48±0.61 µg/ml, P=0.175) compared to placebo group (5.13±0.46 vs. 5.46±0.58 µg/ml, P=0.255) (Fig. 4B). The Δ values both for ICAM-1 (-0.99±6.12 vs. 8.94±4.10 ng/ml, P=0.285) and sVCAM-1 (-0.63±0.46 vs. -0.33±0.28 µg/ml, P=0.927) also showed no difference between the two groups (Fig. 4A and B). Nicorandil may improve the angiokinetic factor of ICAM-1 to hinder any inflammatory reaction during PCI in ACS patients.

**Discussion**

The present study presents for the first time the effects of intra-coronary (IC) bolus injection of nicorandil or placebo on markers of inflammation and oxidative stress in patients with acute coronary syndrome (ACS) undergoing percutaneous coronary interventions (PCIs). In this randomized, single-blinded and placebo-controlled study, it was confirmed in a brief period that percutaneous transluminal coronary angioplasty (PTCA) results in the upregulation of inflammation and oxidative stress. Coronary blood, which has a greater impact on the coronary artery compared with the peripheral blood, was used to evaluate serum as well as plasma levels of inflammatory and oxidative markers to precisely investigate temporal changes during PCI. It was also demonstrated that an IC bolus injection of nicorandil in patients with ACS is safe and significantly reduced soluble CD40 ligand (sCD40L), superoxide dismutase (SOD), malondialdehyde (MDA) and intercellular adhesion molecule-1 (ICAM-1) levels after PCI. However, no changes in other systemic inflammation markers, such as high-sensitivity C-reactive protein (hs-CRP), tumor necrosis factor α (TNFα) and soluble vascular cell adhesion molecule-1 (sVCAM-1) levels were observed in the nicorandil or control groups. This finding could possibly be due to delayed inflammatory responses.
PCI has become a standard revascularization procedure and the main management option for patients with ACS. However, PTCA induces vascular injury and wreaks damage on the endothelium which is associated with pro-inflammatory responses and platelet-activating effects with adhesion (20,21). This damage also elicits an inflammatory response via the activation of platelets and leukocytes (22). PCI-induced vascular injury and ischemia/reperfusion injury was found to result from both intracellular calcium loading due to ATP depletion and intracellular acidification (23), and the occurrence of oxidative stress with the mitochondrial respiratory chain as the main source of reactive oxygen species (ROS) (24).

Nicorandil, as a hybrid nitrate and ATP-sensitive potassium channel opener agent (25), has been shown not only to dilate systemic veins, but also to exert a vaso-dilative effect on arteries, including peripheral arteries (16). Moreover, IC bolus injection of nicorandil is safe and efficacious for myocardial protection, ameliorating fractional flow reserve (FFR) and ischemia/reperfusion injury (26,27). Kawai et al found that intravenous administration of nicorandil could prevent slow coronary flow phenomenon compared with that of 0.9% saline before PCI in patients with ACS or non-ACS (28). Previous research further found that IC bolus injection of nicorandil is safe and effective compared with IC adenosine to induce hyperemia in patients with an angiographically intermediate lesion (27). Lee et al compared IC bolus injection of nicorandil with intravenous infusion of adenosine (26). Different from previous studies, we firstly IC injected the same volume of saline in ACS patients since no placebo effect was observed in those studies which further increased the importance of our study. We also collected coronary blood to perform ELISA to evaluate the levels of inflammatory or oxidative indicators during PCI. A previous study also showed that nicorandil improves microvascular dysfunction as assessed by the index of microcirculatory resistance after PCI (29).

The soluble CD40 ligand (sCD40L), as a member of the TNF family, is expressed widely on monocytes, dendritic cells, endothelial cells and epithelial cells (30). Activation of platelets was shown to result in the release of sCD40L, accelerating the inflammatory process and promoting coagulation (31-34). A previous study showed that sCD40L may be a predictor of coronary artery disease in patients undergoing PCI (35). Thus, we evaluated the level of sCD40L in pre-PCI and post-PCI subgroups in the nicorandil and control groups. Its expression level in intracoronary plasma increased after the stent implantation compared with pre-PCI in the placebo group which is consistent with other trials (35,36). The IC bolus injection of nicorandil also improved the alleviation of sCD40L in the post-PCI subgroup immediately as there were no significant changes compared with the post-PCI and pre-PCI subgroup. Our trial further confirmed that nicorandil has the ability to improve the inflammatory reaction of sCD40L during PCI in ACS patients.
Other inflammation indicators, such as TNFα and hs-CRP, had no significant differences in both nicorandil and placebo groups and their subgroups. Although PCI led to a platelet-activating effect with adhesion and sCD40L participates in the promotion of coagulation, the adhesion molecule sVCAM-1 also exhibited no difference in the different groups although ICAM-1 was clearly induced in the control group after PCI. The reasons for this finding may be as follows. Firstly, Aströmöllson et al demonstrated that TNFα increased during the 24 h of reperfusion (37). Hs-CRP, an acute-phase reactant, increased markedly at 6 h after PCI and reached a peak value at 48 h which indicates that there is a delayed reaction of these inflammatory markers after PCI (37,38). Consequently, the underlying improvement of inflammatory indicators with nicorandil may not react and reflect changes immediately after operation. Secondly, Bayata et al measured VCAM-1 and sICAM-1 serum levels in patients before and after stent implantation and found no difference in VCAM-1 and ICAM-1 levels (39). Similarly, Wexberg et al found increased levels of VCAM-1 after stent implantation (40). Thus, the differences between the above trials and our results may be due to different sourcing of serum or plasma as our samples came from the intracoronary blood drawn by the micro-catheter. Another reason may be a difference in patient ethnicity and the reactivity to PTCA. The small sample size may also be one of the reasons for this difference.

Reactive oxygen species (ROS) are involved in physiological cell regulation and redox signaling (41). SOD, which catalyzes the metabolism of the superoxide anion ($O_2^-$) into hydrogen peroxide and molecular oxygen, is one of the most important antioxidative enzymes. MDA, the major end product of lipid peroxidation, is one of the markers indicating partial levels of oxidative stress. The production of MDA is initiated promptly after the generation of ROS (42). Both molecules directly reflect the expression levels of oxygen-free radicals. Our results showed that both SOD and MDA levels have the tendency to rise but there were no significant differences in both subgroups of the control group. An investigation by Ekele et al indicated that SOD was reduced significantly at 2 h after PTCA (42), which suggests that the consumption of SOD may require a series of processes. However, a study by Demircan et al indicated that SOD was immediately reduced after PCI compared to the pre-PCI subgroup (43). These controversial results require further investigation due to the limited sample sizes employed.

In the present study, it was demonstrated that the SOD levels in the post-PCI subgroup of the nicorandil group significantly increased compared with the pre-PCI subgroup. However, the MDA levels were unexpectedly increased. The ratio of SOD and MDA exhibited no significant change in the post-PCI nicorandil group compared with the control groups. These results showed that nicorandil reduced oxidative stress significantly as a supplement to PCI. The over-production of free radicals at the beginning of the ischemic phase is thought to be due to mitochondrial depolarization. Nicorandil reduces ROS formation by mitochondria by opening the mitochondrial $K_{\text{ATP}}$ (44). Due to its free radical scavenging properties and inhibition of oxidative respiratory bursting of neutrophils (45), IC bolus injection of nicorandil could function by reducing oxidative stress. Furthermore, nicorandil plays a role in ameliorating myocardial damage as well as improving cardiac function and clinical outcomes.

The inflammatory response evoked by vascular damage during angioplasty and the subsequent reaction are thought to be the main contributors to the development of restenosis. Oxidative stress also contributes to clinical outcomes such as adverse cardiac events. IC bolus injection of nicorandil before implantation of the coronary artery stent was associated with better suppression of various inflammatory cytokines, and further reduced the oxidative stress in patients who suffered from ACS undergoing PCI. However, there were no definite clinical benefits detected in the amelioration of adhesion molecules and inflammatory mediators. Our findings suggest that IC bolus injection of nicorandil is a simple, safe, and effective way to reduce the levels of inflammation and oxidation during PCI which may be attributable to improved outcomes for such ACS patients.

There were several limitations to the present study. Firstly, the sample size of this study was small. It was necessary to perform a power analysis before conducting the study. However, no similar data could be found in the published articles which made it difficult for us to assess the sample size. Thus, we enrolled 65 patients to ensure the normality of the data distribution. We surprisingly found significant differences of sCD40L, SOD, MDA and ICAM-1 between the two groups. Then, we performed the power analysis through PASS based on our expression levels of sCD40L, SOD, MDA and ICAM-1. We found that the value of 1-β was beyond 0.8. Secondly, since nicorandil was only used during PCI and only 20 of the enrolled patients took nicorandil regularly post-discharge, no long-term observations of nicorandil was obtained. We followed up the enrolled patients for only one month after PCI. No difference in complications and mortality was found in the two groups after one month. After surgery, the patients recovered well after regular intake of the standard medicine. Finally, according to the new guidelines of ECS for ACS patients, ticagrelor is recommended for Class IA. However, ticagrelor is recommended for Class IB in China at present. According to the 2016 Ticagrelor Expert Consensus in China (46), ticagrelor should be used as early as possible in NSTE-ACS patients who have a moderate or high risk of ischemia and plan for early invasive treatment. We previously used clopidogrel more often during our experimental period according to our previous guidelines. More than 50% patients now use ticagrelor instead of clopidogrel at present. We still need to consider our clinical practice, Chinese national conditions, individual differences and implement specific antithrombotic strategies to choose the antiplatelet drugs. We will also follow up the patients taking nicorandil after PCI for a longer period to explore its long-term effect when we carry out a larger sample study in future studies.

In conclusion, our trial showed that IC bolus injection of nicorandil as part of therapy for patients with ACS undergoing PCI reduced soluble CD40 ligand levels, increased MDA and SOD levels, and decreased ICAM-1. However, none of the other subclinical markers of inflammation were altered after the injection of nicorandil and PCI surgery. These findings within the context of this clinical trial suggest a solitary anti-oxidative and anti-inflammatory effect of nicorandil. A larger sample size and a longer treatment period with nicorandil are needed for further investigation of the underlying and long-term beneficial effects of this therapy.
The prospective, randomized design and use of coronary angiography to diagnose ACS patients are the strengths of this study. The percutaneous transluminal coronary angioplasty (PTCA) showed that a residual stenosis diameter ≥75% of the coronary artery combined with clinical symptoms are the gold standard for identifying ACS patients.

Coronary blood was used to test serum or plasma levels of inflammatory and oxidative markers, which was a better measure of the coronary artery compared with peripheral blood as it precisely reflected the effect on markers after stent deployment.

The limitations of this study included a limited sample size of 65 subjects, inflammatory or oxidative indicators influenced by PTCA and limited time period for extraction of coronary blood during PCA and stent deployment.

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Availability of data and materials

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Authors' contributions

MN, XX and XL conceived, designed and performed the experiments. MN performed the PCI procedures and extracted coronary blood at the appropriate time. XX, JM, SY and LY collected and analyzed the data. All authors contributed to the interpretation of data, drafted or revised the paper. All authors read and approved the manuscript and agree to be accountable for all aspects of the research in ensuring that the accuracy or integrity of any part of the work are appropriately investigated and resolved.

Ethics approval and consent to participate

This study was approved by the Regional Ethics Committee of Shandong University Qilu Hospital, China (Reference no. KYLL-2015266). All members of the operation team and patients were informed of the PCI operation, stent implantation, nicorandil injection and coronary blood extraction. Written consent was obtained before PCI and PTCA. This was approved by the Regional Ethics Committee.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

References

Nicorandil Alleviates Inflammation and Oxidative Stress After Coronary Stent


