Protective effect of hydrogen-rich saline on ischemia/reperfusion in rat skin flap[*]

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Abstract: Objective: Skin damage induced by ischemia/reperfusion (I/R) is a multifactorial process that often occurs in plastic surgery. The mechanisms of I/R injury include hypoxia, inflammation and oxidative damage. Hydrogen (H_2) gas has been reported to alleviate cerebral I/R injury by acting as a free radical scavenger. Here, we assessed the protective effect of hydrogen-rich saline on skin flap I/R injury. Methods: Abdominal skin flaps of rats were elevated and ischemia was induced for 3 h; subsequently, hydrogen-rich saline (HRS) or physiological saline was administered intraperitoneally 10 min before reperfusion. On postoperative d 5, flap survival, blood perfusion, the accumulation of reactive oxygen species and levels of cytokines were evaluated. Histological examinations were performed to assess inflammatory cell infiltration. Results: Skin flap survival and blood flow perfusion were improved by HRS relative to the controls. The production of malondialdehyde (MDA), an indicator of lipid peroxidation, was markedly reduced. A multiplex cytokine assay revealed that HRS reduced the elevation in the levels of inflammatory cytokines, chemokines and growth factors, with the exception of regulated on activation, normal T-cell expressed and secreted (RANTES) growth factor. HRS treatment also reduced inflammatory cell infiltration induced by I/R injury. Conclusion: Our findings suggest that HRS mitigates I/R injury by decreasing inflammation and, therefore, has the potential for application as a therapy for improving skin flap survival.

Key words: Hydrogen-rich saline, Ischemia/reperfusion, Skin flaps, Oxidative damage, Inflammation
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1 Introduction

Skin flap transfer is a routine practice in wound coverage and reconstruction in plastic surgery. Viability is the most important concern, but unfortunately occasional distal necrosis is a fact of life and good planning is not always sufficient to avoid such an outcome. Ischemia/reperfusion (I/R) injury is believed to be a major cause of flap loss. The formation of reactive oxygen species (ROS) during reperfusion triggers lipid peroxidation, protein and nucleic acid damage (Siemionow *et al.*, 2004), and initiates I/R damage (van den Heuvel *et al.*, 2009). Also, inflammation is considered to be a critical element in the pathogenesis of I/R injury. The massive influx of neutrophils results in tissue injury beyond that caused by I/R alone. Numerous substances have been examined in an attempt to attenuate I/R injury in skin or muscle, such as hydrogen sulfide (Henderson *et al.*, 2010), caffeic acid phenethyl ester (Aydogan *et al.*, 2007), and vitamin E (Arato *et al.*, 2010). Most of these studies have shown positive results but lacked independent large-scale follow-up reports.

Ohsawa *et al.* (2007) reported that inhalation of H_2 gas could selectively mitigate •OH damage in a rat model of middle cerebral artery occlusion, generating an antioxidant effect without affecting the signaling of other ROS. Further research demonstrated that the inhalation of H_2 gas produced protective effects on I/R injury in various settings, such as myocardial infarction (Hayashida *et al.*, 2008), intestinal injury (Buchholz *et al.*, 2008), and hepatic injury (Fukuda *et al.*, 2007). Furthermore, H_2 has been reported to downregulate ConA-induced mouse liver inflammation (Kajiya *et al.*, 2009) and to protect mice against multiple organ damage in a zymosan-induced generalized model of inflammation (Xie *et al.*, 2010).

In contrast to other therapeutic but potentially toxic antioxidant gases (nitric oxide, carbon monoxide, hydrogen sulfide, and ozone), hydrogen is less expensive and safer for clinical application (Nakao *et al.*, 2007). However, the clinical application of H_2 gas is not convenient and may be dangerous because H_2 gas is flammable and explosive. H_2 gas-saturated physiological saline, or hydrogen-rich saline, is easy to use and safe to apply. To our knowledge, no published data exist on the protective effect of HRS on skin flap I/R injury. The primary objective of the present study was to evaluate the effect of HRS on the viability of abdominal skin flaps after I/R.

2 Materials and methods

2.1 Animals

All experiments were approved by the Committee on Animal Rights Protection of Peking Union Medical College Hospital and were performed in accordance with the National Institutes of Health (NIH) guidelines for the care and use of laboratory animals.

Adult male Sprague-Dawley (SD) rats weighing 270–320 g were used in this study. The rats were housed under standard conditions at 22 °C to 25 °C with a 12 h light-dark cycle and were fed a normal diet with water provided ad lib pre- and postoperatively.

2.2 HRS production and distribution studies

HRS was made by dissolving hydrogen in normal saline (6 h) under high pressure (0.4 M) until supersaturated. The hydrogen content was confirmed by the method described by Ohsawa *et al.* (2007). Fresh HRS was produced weekly to ensure supersaturation. The H_2 concentration was maintained above 0.6 mM. Any remaining solution was discarded after the package was opened or damaged.

The rats were anesthetized with an intraperitoneal (i.p.) injection of pentobarbital at a dose of 40 mg/kg body weight (BW) and placed in a supine position. The rats were also i.p. injected with 5 ml/kg BW HRS, 10 ml/kg BW HRS, or the saline control. The H_2 concentration in the skin flap tissue was measured using a needle-type H_2 sensor (Unisense, Denmark).
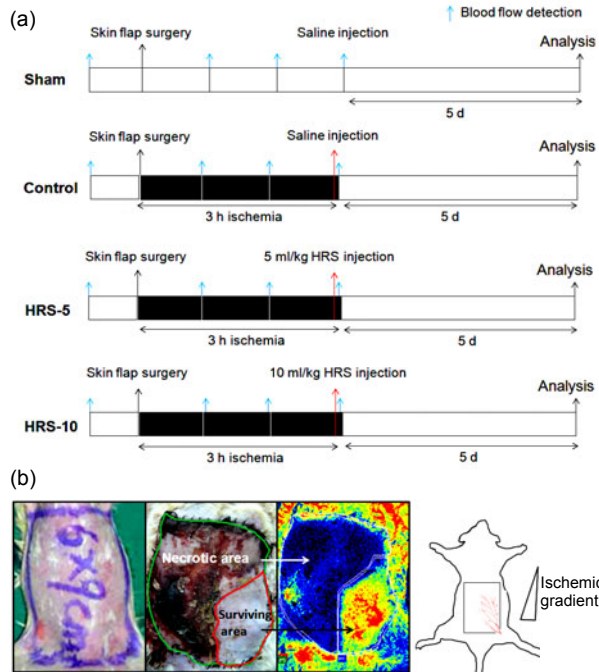
2.3 HRS production and distribution studies

Sixty male SD rats were divided randomly into four groups consisting of 15 animals per group: (1) a sham-operated group: no I/R, HRS, or normal saline injection. (2) a control group: 3 h ischemia induced by clamping the right pedicle followed by an i.p. injection of 5 ml/kg normal saline 10 min before reperfusion; and (3) two hydrogen-rich saline treated groups: 3 h ischemia induced and followed by an i.p. injection of 5 ml/kg HRS (HRS-5 group) or 10 ml/kg HRS (HRS-10 group), 10 min prior to reperfusion (Fig. 1a).

2.4 Epigastric skin flap preparation and survival

An extended epigastric adipocutaneous flap (6×9 cm) was raised in each animal (Küntscher *et al.*, 2007). The left superficial epigastric artery and vein were ligated so that the blood was supplied only by the right pedicle. A microvascular clamp was used to occlude the artery and vein, inducing skin flap ischemia (3 h). The flap was resutured, and a gap was left for the removal of the clamp. To prevent neovascularization in the wound bed, a silicone sheet was placed between the flap and the abdominal muscle bed. At the end of the ischemic period, the clamp was removed, and heparin (50000 U/L in 0.5 ml saline) was used to avoid thrombus formation.

Skin flap survival was evaluated by the general observation of survival and necrotic phenomena and subsequently confirmed by Laser Speckle Contrast Analysis (LASCA) cameras (Perimed AB, Sweden). The surviving and necrotic areas were measured using Acrobat 8.0 software. The percentage of flap survival was defined as the ratio of the surviving area to the original flap area (Fig. 1b).

Fig. 1 Schematic illustration of the experimental protocol used to determine the effect of HRS on I/R skin flaps
(a) Experimental protocol: sixty SD rats were divided into four groups as described in the Materials and methods section. Occlusion (3 h) induced ischemia, and HRS or saline was injected. Blue arrows indicate blood flow detection, and red arrows indicate reperfusion time points. (b) A rectangular 6×9 cm skin flap was raised in each animal and was supplied by the left superficial epigastric artery. Surgery led to ischemia and subsequent necrosis in the distal part of the ski flap. The percentage of tissue survival was quantified by planimetry, calculating the area of survival (red area) divided by the area of the skin flap (green area)

2.5 Flap perfusion

Before scanning, the rats were secured onto the operative bed so that the whole flap, including part of the normal abdominal skin, was exposed. The PeriScan PSI system was positioned above the rats so that an 11×7.5 cm area was imaged. The image acquisition rate was 3 s^{-1} and lasted for about 3 min. The ambient temperature was maintained between 22 and 25 °C during this process.

On the operative day, blood flow perfusion was detected before surgery and at 1, 2, and 3 h after surgery, denoted Hour 0, Hour 1, Hour 2, and Hour 3, respectively. Perfusion of the necrotic and survival areas was analyzed on the fifth postoperative day. Vascular flow was measured using PUs (Perfusion Units).

2.6 Hematoxylin and eosin (H&E) staining

On the fifth postoperative day, three pieces of the flap tissue (each about 1 cm^2 in size) were taken from the surviving areas of the flap. The harvesting locations were proximal, middle and distal along the vascular axis. Samples were sectioned into pieces and fixed with 4% paraformaldehyde in 0.1 mol/L phosphate buffer. They were then embedded in paraffin, mounted onto slides, and stained with hematoxylin and eosin.

2.7 Malondialdehyde (MDA) measurement

Briefly, 25 mg of frozen flap tissue, cut into pieces on ice, were homogenized in RIPA buffer using a manual glass homogenizer, and centrifuged for 10 min. The supernatant was used for analysis. The MDA content of homogenates was determined spectrophotometrically by measuring the presence of thiobarbituric acid reactive substances (Cayman, USA). Results are expressed as [MDA] in nmol/mg of protein.

2.8 Multiplex cytokine assay

A rat 23-plex kit (Bio-rad Laboratories, Hercules, CA) was used to determine the concentrations of 23 cytokines simultaneously in the surviving flap tissues, including interleukin (IL)-1α, IL-1β, IL-2, IL-4, IL-5, IL-6, IL-7, IL-10, IL-12, IL-13, IL-17, IL-18, tumor necrotic factor (TNF)-α, macrophage inflammatory protein (MIP)-1α, MIP-3α, interferon (IFN)-γ, erythropoietin (EPO), macrophage colony-stimulating factor (M-CSF), granulocyte-macrophage colony-stimulating factor (GM-CSF), granulocyte colony-stimulating factor (G-CSF), vascular endothelial growth factor (VEGF), regulated on activation, normal T-cell Expressed and Secreted (RANTES) growth factor, and growth-regulated oncogene /KC (GRO/KC). The samples (100 mg per sample tissue) were homogenized using a cell lysis kit (Bio-Rad Laboratories, Hercules, CA), and the protein concentrations were measured by the Bradford assay (Tiangen, Beijing, China).

In brief, the lysates were diluted to 900 μg/ml, and 50 μl of each diluted sample was incubated with antibody-coupled beads. The complexes were washed, incubated with a biotinylated detection antibody and then incubated with streptavidin-phycoerythrin prior

to assessing the cytokine concentration titers. Assay standards resuspended in standard diluents were used to plot the standard curves. Cytokine concentrations were determined by comparison against a set of standards. The data were automatically processed and analyzed by Bio-Plex Manager Software 4.0.

2.3 Statistical analysis

All data are reported as the mean±standard error of the mean (SEM). The differences between the levels of cytokines in each group were determined via one-way analysis of variance (ANOVA), and the Student's t-test was used to analyze the differences between the surviving and original flap areas. A 2-tailed probability level of $P < 0.05$ was used to indicate statistical significance. Non-significant results are reported as 'ns'.
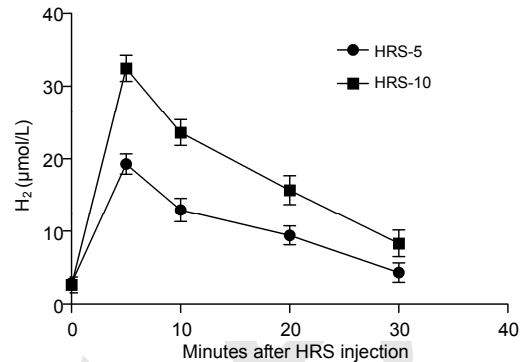
3 Results

3.1 Intraperitoneal injection of HRS significantly increased the H_2 concentration in the flap

A linear correlation was found between the current value of the H_2 microelectrode and the hydrogen concentration (H_2 concentration: 0–50 μmol/L, R^2=0.9993; data not shown). H_2 concentrations peaked at about 5 min after the HRS injection (Fig. 2). HRS treatment at 5 ml/kg increased the H_2 concentration to (19.14±1.41) μmol/L, while 10 ml/kg HRS increased the H_2 concentration to (32.18±0.56) μmol/L. These results suggest that i.p. administration of HRS may efficiently deliver H_2 into skin flaps. Furthermore, we found no significant difference in the pH values between HRS and normal saline (HRS, 7.32±0.02; saline, 7.34±0.03).

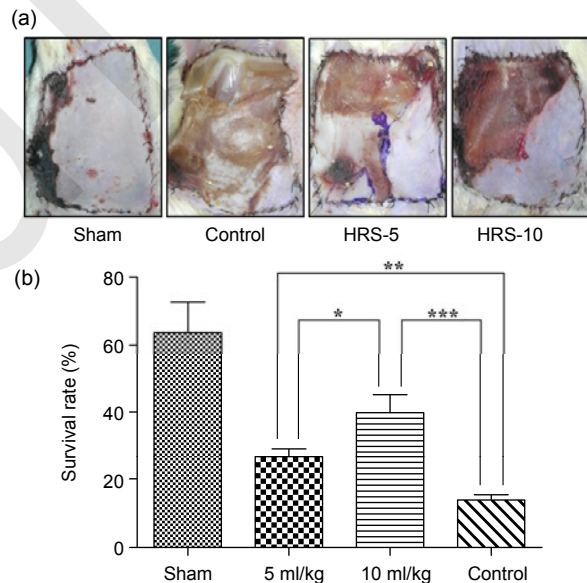### 3.2 Flap survival was improved by HRS treatment

On the fifth postoperative day, necrotic tissues were observed and presented as grey, brown, or black areas with little elasticity. In contrast, the tissue in surviving areas maintained normal elasticity and the skin color was pink and white (Fig. 3a). The quantitative analysis showed that survival in the control group accounted for (13.92±1.53)% of the total flap area, whereas it represented (26.52±2.60)% in the HRS-5 group ($P < 0.01$ vs. control) and (39.85±5.32)% in the HRS-10 group ($P < 0.001$ vs. control) (Fig. 3b).

The control group flap showed a variable degree of full-thickness necrosis and petechiae. These parameters were significantly diminished in the HRS groups, suggesting a protective effect of HRS treatment on skin flap necrosis.

Fig. 2 Intraperitoneal injection of HRS increases H_2 concentration in the skin flaps.
5 ml/kg or 10 ml/kg HRS was injected. The rats were anesthetized with pentobarbital (40 mg/kg BW). A H_2 microelectrode (diam. 50 μm) was inserted into the skin flaps. H_2 concentration peaked at about 5 min post-injection

Fig. 3 HRS significantly increased the survival rate of skin flaps on postoperative Day 5
(a) Rats were anesthetized with i.p. pentobarbital injection and secured to the operative bed. The image was captured with a digital camera. Obvious necrosis was observed in the control group, whereas HRS significantly improved the survival rate in both the HRS-5 and HRS-10 groups. (b) Flap survival was assessed using the methods described in Fig. 1b. Survival area percentages were (13.92±1.53)% of the total flap area in the control group compared to (26.52±2.60)% in the HRS-5 group and 39.85±5.32% in the HRS-10 group. * $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$

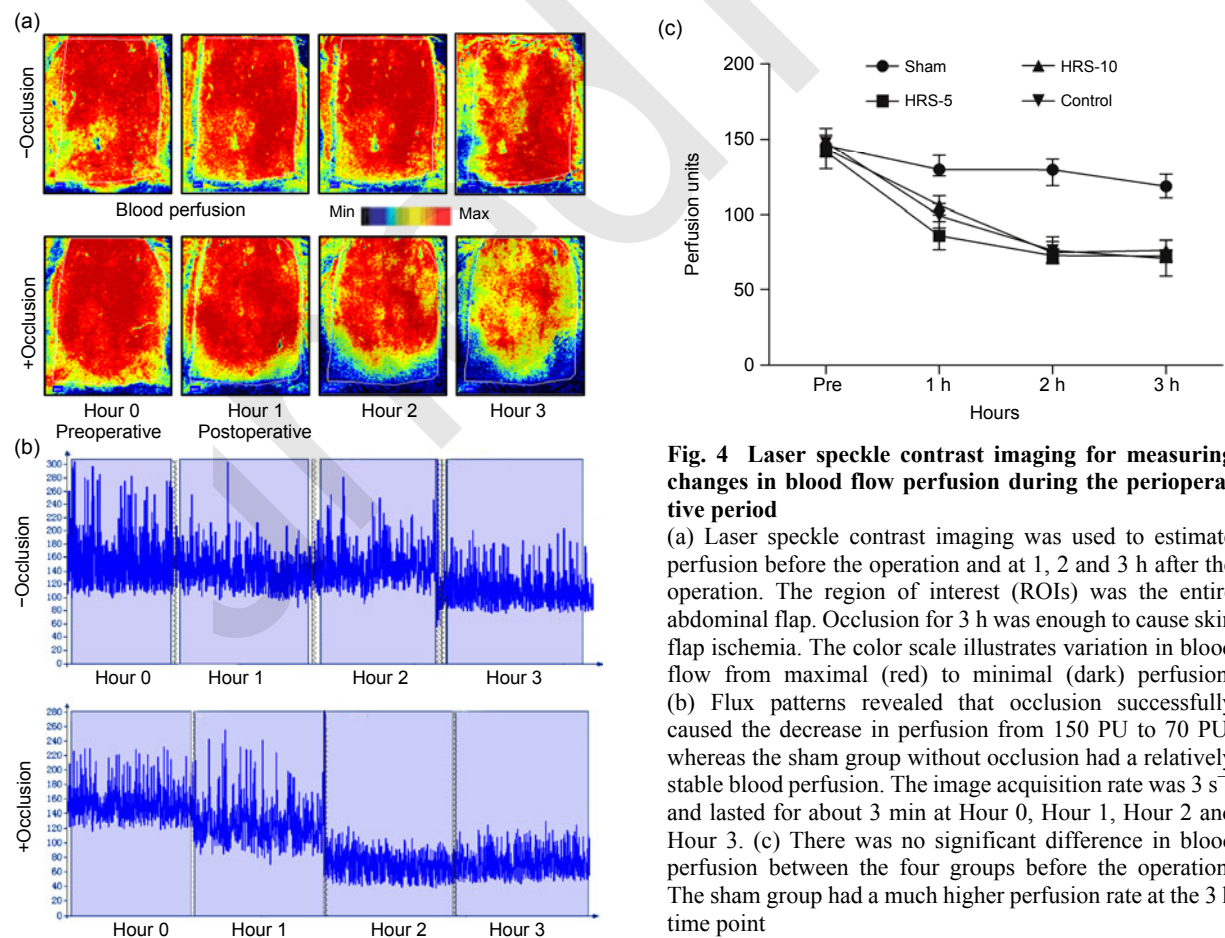## 3.3 Flap perfusion was ameliorated by HRS treatment

The evaluation of skin flap perfusion was accomplished as described in the Materials and methods section. As we expected, occlusion (3 h) significantly decreased blood flow (Figs. 4a and 4b). There were no significant differences in the decrease in blood perfusion among the three operative groups (HRS-5, HRS-10 and control group). In contrast, the non-occluded sham group showed relatively stable and much better perfusion (Fig. 4c). These results indicate that a 3-h occlusion was enough to induce flap ischemia.

Because the flap was axial and centered on the right superficial epigastric artery and vein, the restoration of blood flow following severe ischemia began from the proximal portion of the flap. On postoperative Day 5, perfusion was only (38.98±3.81) PU in the control group. A significantly higher blood perfusion was measured in the HRS groups compared with the controls (Fig. 5a). The quantification of blood flow

was (71.32±12.88) PU in the HRS-5 group and (81.79±13.56) PU in the HRS-10 group ($P<0.05$ vs. control). There was no significant difference in the blood flow levels between the HRS-5 and HRS-10 groups ($t=3.17$, $P>0.05$) (Fig. 5b). These data suggest that HRS treatment significantly improves blood flow in skin flaps during reperfusion.
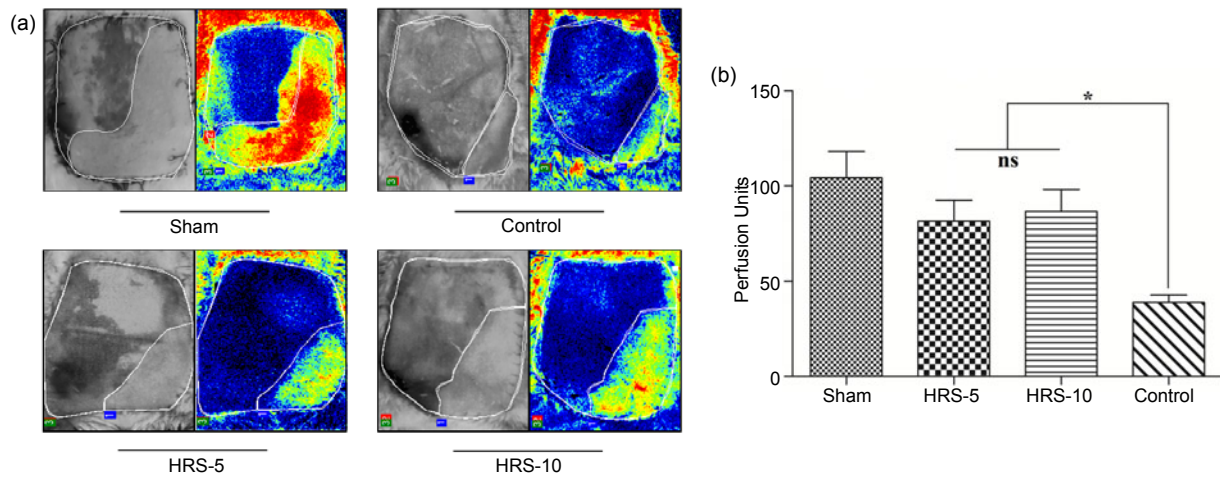
3.4 Histological analysis

Histological analysis of H&E stained tissues was performed to qualify ischemic lesions. As we expected, inflammation was progressively more severe moving from the proximal to distal portion of the tissue, as denoted by an increase in the number of inflammatory cells. No significant infiltration was observed in the sham-operated animals. In contrast, a significant number of infiltrated cells were observed in the skin flaps of the untreated rats in the control group. Thus, treatment with HRS decreased this infiltration, suggesting that HRS attenuated activation of the inflammatory response (Fig. 6).

Fig. 4 Laser speckle contrast imaging for measuring changes in blood flow perfusion during the perioperative period
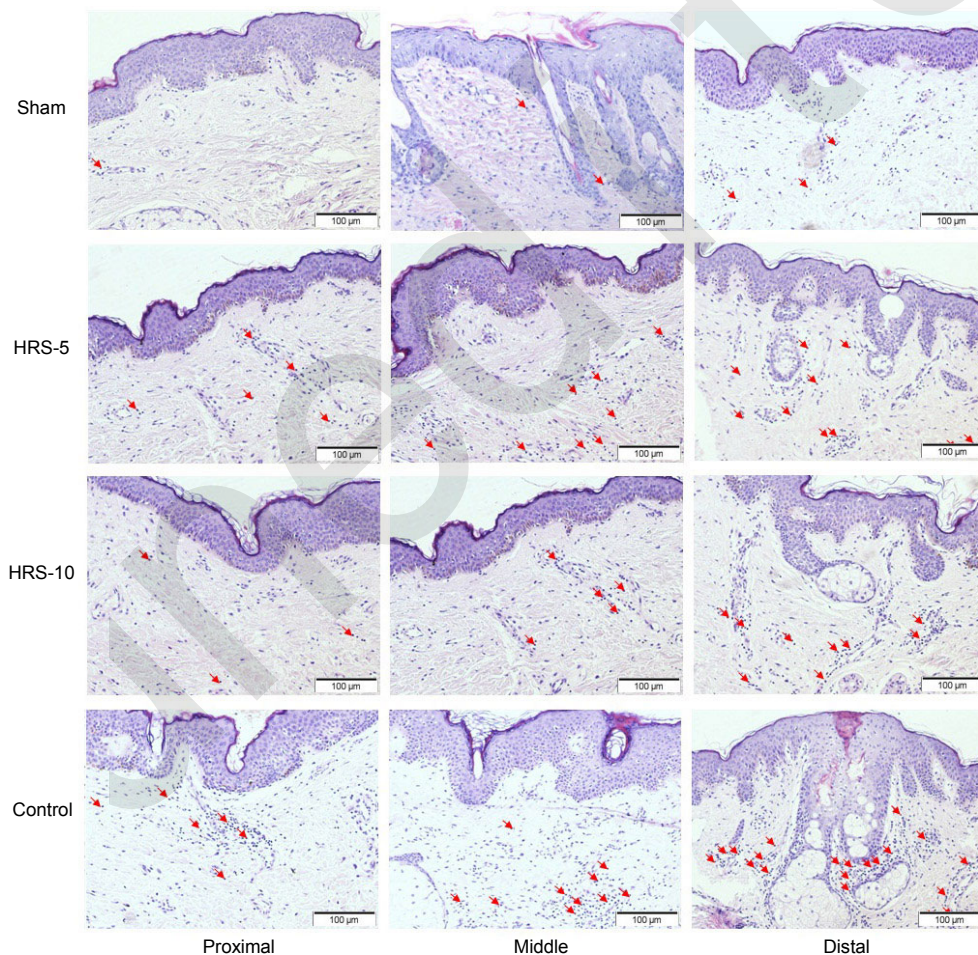
(a) Laser speckle contrast imaging was used to estimate perfusion before the operation and at 1, 2 and 3 h after the operation. The region of interest (ROIs) was the entire abdominal flap. Occlusion for 3 h was enough to cause skin flap ischemia. The color scale illustrates variation in blood flow from maximal (red) to minimal (dark) perfusion. (b) Flux patterns revealed that occlusion successfully caused the decrease in perfusion from 150 PU to 70 PU, whereas the sham group without occlusion had a relatively stable blood perfusion. The image acquisition rate was 3 s^{-1} and lasted for about 3 min at Hour 0, Hour 1, Hour 2 and Hour 3. (c) There was no significant difference in blood perfusion between the four groups before the operation. The sham group had a much higher perfusion rate at the 3 h time point

Fig. 5 Blood flow perfusion analysis using LACSA on postoperative d 5

(a) The blood flow perfusion in the surviving and necrotic areas was detected. The images demonstrate that HRS ameliorated skin blood flow perfusion in both HRS treatment groups. (b) Quantitative analysis of blood flow on postoperative d 5. HRS groups had higher blood perfusion compared with the controls. Blood perfusion is expressed as the mean±SEM. $^*P<0.05$
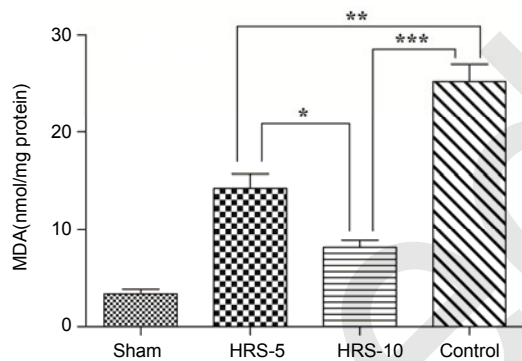

Fig. 6 Photomicrographs of the middle area of skin flaps in different groups

The number of infiltrated cells increased from the proximal to the distal portions of the skin flaps. Normal skin tissue was observed, and no obvious inflammation was observed in the sham group. We observed significant cellular infiltration in the control group, suggesting higher activation of the inflammatory process. In contrast, HSR-5 and HSR-10 treatments significantly decreased cellular infiltration. The red arrows indicate inflammatory cells. (100 X)

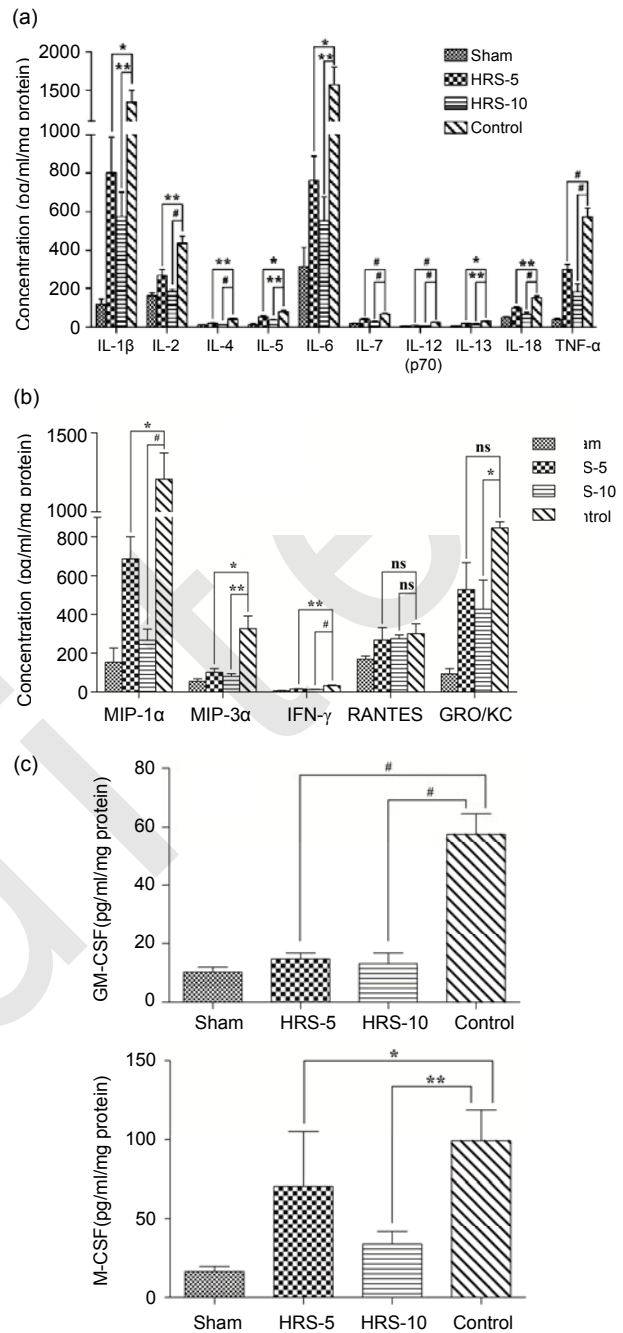## 3.5 HRS treatment inhibited the elevation of MDA

As a measurement of lipid peroxidation, MDA levels in the proximal, middle and distal portions of the skin flap were measured. We found that the MDA levels increased from the proximal portion to the distal portion, but there were no significant differences between the three parts (data no shown). The results are expressed as the average of the three parts and correspond to samples obtained five days after surgery. The rats subjected to skin flap I/R exhibited an increase in MDA levels compared with the sham-operated rats on the fifth postoperative day (Fig. 7). MDA formation averaged ((25.32 ± 1.79) nmol/mg of protein). Treatment with HRS resulted in a marked reduction in MDA levels in the HRS-5 group ((14.27 ± 1.43) nmol/mg of protein) and in the HRS-10 group ((8.17 ± 0.72) nmol/mg of protein). These results suggest that HRS treatment significantly attenuated lipid peroxidation in the I/R skin flaps.

Fig. 7 HRS decreased production of MDA in the flap
The control group exhibited a significantly higher MDA level, whereas HRS decreased the production of MDA. The results are expressed as the mean±SEM. $^*P<0.05$, $^{**}P<0.01$, $^{***}P<0.001$

3.6 Cytokine evaluation

Cytokines in the proximal, middle and distal portions of surviving tissue were detected, and the data are reported as the average of the three parts. We successfully detected 18 of all 23 cytokines. The levels of 18 cytokines were higher in the control group than in the sham-operated group. HRS treatment evidently prevented the upregulation of inflammatory mediators, including IL-1β, IL-2, IL-4, IL-5, IL-6, IL-7, IL-12, IL-13, IL-18, and TNF-α (Fig. 8a).

Fig. 8 Cytokine profiles on skin flaps
Tissues were harvested at postoperative day 5 from the proximal, middle and distal points along the vessel axis. Cytokine production was assessed by multiplex analysis of tissue homogenates. (a) HRS administrated 10 min before reperfusion markedly reduced the upregulation of all inflammatory cytokines. (b) Chemokines were reduced by HRS treatment, with the exception of RANTES. GRO/KC was markedly reduced in the HRS-10 group. (c) HRS treatments decreased the levels of GM-CSF and M-CSF in the tissues. Cytokine changes indicated the alleviation of inflammation. $^*P<0.05$, $^{**}P<0.01$, $^#P<0.001$

I/R injury also induced the elevation of the concentrations of the detected chemokines (Fig. 8b). MIP-1α, MIP-3α, and IFN-γ levels were reduced by HRS in both the HRS-5 and HRS-10 groups. GRO/KC, a potent neutrophil chemoattractant, was reduced by higher-dose HRS administration. However, there was no significant difference in the RANTES level between the HRS group and the control group.

GM-CSF is a cytokine that functions as a white blood cell growth factor. M-CSF is a hematopoietic growth factor that is involved in the proliferation and differentiation of monocytes and macrophages. HRS treatment reduced the up-regulation of GM-CSF and M-CSF compared to the controls (Fig. 8c).

4 Discussion

In this study, we report the protective effect of HRS on skin flaps after 3 h of ischemia. Our data demonstrate that intraperitoneal HRS administration improved skin flap survival, ameliorated microvascular blood flow disturbances, and reduced MDA production. HRS also alleviated inflammation-related events by reducing leukocyte infiltration and pro-inflammatory cytokine production. Histology indicated that the recruitment of leukocytes was reduced, and sixteen measured inflammatory-relevant cytokines were downregulated by HRS treatment. Taken together, these results indicate that HRS may be a novel antioxidant and anti-inflammatory agent with a promising role in protecting skin flaps from I/R injuries.

Flap complications commonly occur in skin flap transfer procedures for wound coverage and reconstruction in plastic surgery. Although many strategies have been investigated to prevent flap morbidity, including pharmaceutical agents, preconditioning (Dacho *et al.*, 2009; Küntscher *et al.*, 2002), and postconditioning (Yan *et al.*, 2010; Moon *et al.*, 2008), the total loss of microsurgically transferred flaps is about 1-5% in experienced hands (Harder *et al.*, 2010). The partial flap necrosis rate is 7-20% of free flaps and 20-30% of pedicled flaps. I/R injuries are an important reason for flap necrosis and failure.

During reperfusion many ROS evolve in a short period of time, thereby flooding the antioxidant system. •OH is the strongest of the oxidant species, and mammalian species lack endogenous detoxification systems to neutralize it. Therefore, the therapeutic targeting of •OH could be critical for the amelioration of oxidative injury. Free-radical scavengers can convert toxic ROS into common and relatively harmless end products. Many antioxidants, such as superoxide dismutase (Manson *et al.*, 1983), grape seed proanthocyanidin extract (Karaaslan *et al.*, 2010), and melatonin (Gurlek *et al.*, 1996), have been used to limited tissue injury. However, strategies that decrease oxidative status intensively may produce unwanted side effects because at low levels ROS function as signaling molecules to regulate apoptosis, cell proliferation and differentiation.

H_2 was reported as a new antioxidant and a selective scavenger of •OH effective in various organs (Zheng *et al.*, 2009; Sun *et al.*, 2011; Nagata *et al.*, 2009). H2 can convert hydroxyl into water (H_2+•OH →H_2O_2+•H). Because H_2 is electronically neutral and a small molecule, it should easily penetrate the cellular and intracellular membranes that are normally barriers preventing water-soluble antioxidants from entering cells and organelles, such as the mitochondria, a major source of ROS production. We monitored the concentration of H_2 in skin flaps using real-time dynamic methods and found that the skin flap H_2 concentration reached a peak at about 5 min after HRS intraperitoneal injection. MDA production during free radical attack on membrane lipoproteins and polyunsaturated fatty acids is an indicator of lipid peroxidation. In this study, the notable increase in MDA levels in the control group confirmed the oxidative damage in the skin flap. In contrast, HRS treatments markedly reduced MDA production; thus, we conclude that the intraperitoneal administration of HRS may prevent damage associated with ROS.

Another important factor contributing to flap damage is the release of inflammatory mediators during reperfusion. ROS can activate TNF-α and IL-1β expression by upregulating the NF-kB signaling pathway. Subsequently, pro-inflammatory cytokines (TNF-α, IL-1, IL-6), chemokines (MIP-1α), and growth factors (GM-CSF, M-CSF) cause macrophage maturation, which further activates the immune/inflammatory cascade and results in the secretion of more inflammatory cytokines. On postoperative d 5, inflammation remained severe in the controls

as manifested by the higher expression of cytokines and increased inflammatory cell infiltration relative to the sham-operated group. HRS administration attenuated IR-induced expression of pro-inflammatory cytokines, growth factors, and cytokines (IL-2, IL-3, IL-4, etc.). Histological examination also demonstrated clearly that HRS injection prevented structural skin damage with mild inflammatory accumulation. Given the complexity of cross-reactions between ROS and pro-inflammatory mediators, the H_2-mediated suppression of inflammation induced by I/R injury may also involve antioxidant effects of H_2.

Many materials are known to increase flap viability, but the majority of these agents with proven efficacy require systemic and relatively high dosages, which may cause system side effects. H_2 can reach relatively high concentrations quickly, and excessive H_2 can be eliminated from the body via breathing, causing no side effects (Chen *et al.*, 2011). In addition, therapeutic H_2 effects on humans have been reported for diabetes mellitus type 2 (Imai *et al.*, 2008), hemodialysis (Nakayama *et al.*, 2010), inflammatory myopathies (Ito *et al.*, 2011), radiotherapy for liver cancer (Kang *et al.*, 2011), and acute erythymatous skin diseases (Ono *et al.*, 2012). H_2 can be administered through intravenous injection or drink. Potent antioxidant activity, easy application, lack of significant toxicological effects and cost-effectiveness make HRS an ideal candidate for flap survival in the clinical setting.

5 Conclusions

HRS increased the surviving areas of rat epigastric adipocutaneous flaps and decreased oxidative stress and inflammation in these tissues. Thus, the results of the current study validate the therapeutic potential of HRS by demonstrating that HRS injection 10 min before reperfusion mitigates I/R injury.

Compliance with ethics guidelines

Ling Zhao, Youbin Wang, Shirui Qin, Xuemei Ma, Xuejun Sun, Minglian Wang, and Rugang Zhong declare that they have no conflict of interest.

All institutional and national guidelines for the care and use of laboratory animals were followed.

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