Application of pulsed arterial resuscitation in a rabbit model of hemorrhagic shock

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ABSTRACT

BACKGROUND: Hemorrhagic shock is characterized by tissue hypoperfusion caused by a sharp reduction in the effective circulating volume of blood. The key to successful resuscitation lies in eliminating the shock as soon as possible while simultaneously restoring blood perfusion to vital organs. We present the applicability of pulsed arterial blood reinfusion for resuscitation of hemorrhagic shock.

METHODS: Sixty rabbits were randomly assigned to resuscitation and control groups. A rabbit hemorrhagic shock model was developed by bloodletting from the carotid artery. The dynamic changes in blood pressure, urine output, blood lactate, and other indicators were measured.

RESULTS: Compared with the control group, the mean arterial pressure (MAP), pulse pressure, and urine output were significantly higher in the resuscitation group at 60 min (MAP: 83.67±3.90 vs. 38.19±3.50 mmHg, p<0.001; pulse difference: 16.46±2.21 vs. 10.27±2.99 mmHg, p<0.001; urine output: 3.68±0.74 vs. 0.10±0.05 mL·kg⁻¹·min⁻¹, p<0.001), whereas the serum lactate level was significantly lower (3.82±0.50 vs. 6.49±0.61 mmol/L, p<0.001). In addition, the resuscitation group had a significantly higher lactate clearance rate (30 min: 0.26%±0.11% vs. 0.25%±0.14%, p<0.001; 60 min: 0.30%±0.09% vs. 0.67%±0.26%, p<0.001) than the control group.

CONCLUSION: Pulsed arterial resuscitation might be useful for emergency treatment of hemorrhagic shock.

Keywords: Artery; hemorrhagic shock; pulse; resuscitation.
the bilateral common carotid arteries, if the liquid is infused through one side of the common carotid artery, the infusion will not cause reduction in blood flow to the brain because of the compensation from the other side. Thus, arterial resuscitation may represent an effective option for critically ill patients in whom conventional intravenous resuscitation is not suitable. In this study, we aimed to explore the applicability of pulsed arterial blood reinfusion for the resuscitation of hemorrhagic shock using a rabbit model of hemorrhagic shock. Furthermore, we hypothesized that pulsed arterial reinfusion would be more effective than the traditional liberal fluid resuscitation.

MATERIALS AND METHODS

Animal Grouping and Preparation

Sixty male or female (non-pregnant) rabbits, with a body weight of 1.5–2.5 kg, were provided by the Animal Center of the Southeast University School of Medicine (Animal Certificate of Conformity: SCXK [Su] 2012-0003). The rabbits were 1:1 randomly assigned using a random number table to the resuscitation or control group, with 30 rabbits in each group. After they were weighed, the rabbits were anesthetized by injecting 20% urethane solution (5 mL/kg) through the ear vein and were fixed on the bench. Subsequently, the rabbits were subjected to a midline neck incision. The common carotid arteries were isolated and catheterized to collect blood samples with heparinization. The end of the catheter was connected to a biological signal acquisition system through a sterilized three-way connector (i.e., sterilized T-branch three-way pipe) to trace the arterial blood pressure, electrocardiography parameters, and other vital indicators.[9] The bladder was exposed and catheterized to collect urine specimens. The end of the catheter was connected to a biological signal acquisition system to monitor urine output. The research protocol was approved by the Institutional Review Board and the ethical committees of the Clinical Medical College of Hangzhou Normal University.

Preparation of the Hemorrhagic Shock Model

The model was established using the described method with slight modifications.[4,17] After catheterization was completed and stabilized for 10 min, the blood pressure and heart rate were recorded, and blood samples were obtained as the baseline parameters. Next, blood was withdrawn at 2 mL/min from the common carotid artery through a 50-mL syringe. The blood received an immediate anti-coagulant treatment with pH control and was stored for the subsequent blood re-infusion. Within 30 min after bloodletting, the mean arterial pressure (MAP) had decreased to 50% of the baseline value, which was maintained for 30 min.

Pulsed Arterial Blood Reinfusion Resuscitation

The resuscitation began after the shock had been stable for 30 min. Within 5 s, rabbits in the resuscitation group were re-infused with 5 mL of blood from the carotid artery, followed by 5 mL of saline (using the same lavage tube). This procedure was repeated every 5 min. Rabbits in the control group were re-infused into the venous system with 5 mL of blood from the ear vein at a constant speed, followed by 5 mL of saline (using the same lavage tube). This procedure was repeated every 5 min. The volume of the re-infused blood was 40% of the total blood volume removed.

Monitoring of indicators

Animal systolic blood pressure, diastolic blood pressure, MAP, pulse pressure changes, and urine output were monitored throughout the experiment, and the changes in lip color were continuously observed. At 0, 30, 60 (start of resuscitation), and 90 min after the hemorrhagic shock model was established, blood was withdrawn to measure the lactate level (kit provided by Nanjing Jiancheng Bioengineering Research Institute, Jiangsu, China).

Statistical analysis

All data were analyzed using SPSS 19.0 software (SPSS Inc., Chicago, IL). Quantitative data are presented as mean±standard deviation (x±s) and were analyzed using non-paired t-tests. Qualitative data were analyzed using chi-squared tests. For all analyses, the level of significance was set at p<0.05.

RESULTS

General Condition

At 10 min after bloodletting, the animals began to show cyanosis of the lips, which gradually intensified during the shock process. At the beginning of the resuscitation, there were no significant changes in the lip color. However, when the re-infused blood volume amounted to approximately 30% of the total blood loss, the cyanosis on the lips of the rabbits in the resuscitation group began to subside. At the late stage of the resuscitation, the cyanosis had almost disappeared. In the control group, there were no significant changes in the cyanosis, with the rabbit lips exhibiting a dark purple color at the late stage of the resuscitation.

Mortality Rates

No experimental animals died during the surgical preparation stage or the 30-min shock period. At 60 min, there were no deaths in the resuscitation group, whereas there were 9 deaths in the control group; however, there was no statistically significant difference in the mortality rate between the two groups at this time (p>0.05). At 90 min, there were still no deaths in the resuscitation group, whereas there were 18 deaths in the control group (p<0.01; Table 1).

Comparison of Blood Pressure

There were no significant changes in the blood pressure at
0 and 30 min after the hemorrhagic shock model was established (p>0.05 for both). MAP and pulse pressure of the rabbits in the resuscitation group were significantly increased at 60 min than at 30 min (p<0.05 for both), whereas in the control group, the blood pressure was constantly decreasing. From the beginning of the resuscitation, MAP and pulse pressure of rabbits in the resuscitation group were significantly higher than those of rabbits in the control group at all time points (p<0.01 for all; Table 2).

### Comparison of Urine Output

In both groups, there were no significant differences in the urine output between 0 and 30 min after the hemorrhagic shock model was established (p>0.05 for both). At 60 and 90 min, the urine output of the rabbits in the resuscitation group was significantly increased when compared to that at 30 min, although it was lower than that at 0 min (p<0.05 for all). Conversely, in the control group, the urine output decreased constantly. From the beginning of the resuscitation, the urine output of the rabbits in the resuscitation group was significantly higher than that of the rabbits in the control group at all time points (p<0.01 for all; Table 3).

### Comparison of Blood Lactate Levels

In both groups, there were no significant differences in the blood lactate levels between 0 and 30 min after the hemorrhagic shock model was established (p>0.05 for both). At 60 and 90 min, the lactate level of the rabbits in the resuscitation group was significantly decreased compared to that at 30 min, although it was higher than that at 0 min (p<0.01 for all). Conversely, in the control group, the lactate level decreased constantly. From the beginning of the resuscitation, the lactate level of the rabbits in the resuscitation group was significantly lower than that of the rabbits in the control group at all time points (p<0.01 for all; Table 4).

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**Table 1.** Comparison of the mortality rates between the two groups treated with different reinfusion methods at different time points after establishing the rabbit model of hemorrhagic shock

<table>
<thead>
<tr>
<th>Group</th>
<th>No. of animals</th>
<th>0 min</th>
<th>30 min</th>
<th>60 min</th>
<th>90 min</th>
</tr>
</thead>
<tbody>
<tr>
<td>Resuscitation</td>
<td>30</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>0 (0)</td>
</tr>
<tr>
<td>Control</td>
<td>30</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>30 (9)</td>
<td>60 (18)</td>
</tr>
</tbody>
</table>

χ² value
P value

0.060 0.003

The resuscitation group received pulsed carotid artery reinfusion, and the control group received pulsed intravenous reinfusion. The resuscitation began at 60 min. Non-applicable values are left as blanks in this table.

**Table 2.** Comparison of the dynamic changes in blood pressure between the two groups treated with different reinfusion methods at different time points after establishing the rabbit model of hemorrhagic shock (Mean±SD)

<table>
<thead>
<tr>
<th>Group</th>
<th>Mean arterial pressure (mmHg [no. of animals])</th>
<th>Pulse pressure (mmHg [no. of animals])</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>0 min</td>
<td>30 min</td>
</tr>
<tr>
<td>Resuscitation</td>
<td>103.15±1.54 (30)</td>
<td>50.41±1.89 (30)</td>
</tr>
<tr>
<td>Control</td>
<td>102.65±1.87 (30)</td>
<td>49.47±3.71 (30)</td>
</tr>
<tr>
<td>t value</td>
<td>0.206</td>
<td>0.715</td>
</tr>
<tr>
<td>P value</td>
<td>0.839</td>
<td>0.484</td>
</tr>
</tbody>
</table>

The resuscitation group received pulsed carotid artery reinfusion, and the control group received pulsed intravenous reinfusion. Resuscitation began at 60 min. The value at 60 min was compared with that at 30 min within each group (p<0.05 for all). The number of animals is recorded within parentheses. *1 mmHg = 0.133 kPa.

**Table 3.** Comparison of the dynamic changes in urine output between the two groups treated with different reinfusion methods at different time points after establishing the rabbit model of hemorrhagic shock (Mean±SD)

<table>
<thead>
<tr>
<th>Group</th>
<th>Urine output (mL·kg⁻¹·min⁻¹, Mean±SD [no. of animals])</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>0 min</td>
</tr>
<tr>
<td>Resuscitation</td>
<td>23.67±2.891 (30)</td>
</tr>
<tr>
<td>Control</td>
<td>24.78±2.604 (30)</td>
</tr>
<tr>
<td>t value</td>
<td>-0.904</td>
</tr>
<tr>
<td>p value</td>
<td>0.378</td>
</tr>
</tbody>
</table>

The resuscitation group received pulsed carotid artery reinfusion, and the control group received pulsed intravenous reinfusion. Resuscitation began at 60 min. The value at 60 min was compared with that at 0 min (p<0.05) and at 30 min within each group (p<0.05). The number of animals is recorded within parentheses.
rhagic shock model was established (p>0.05 for both). From the beginning of the resuscitation, the blood lactate level of the rabbits in the resuscitation group significantly decreased, whereas in the control group, the blood lactate level continued to increase. At the beginning of the resuscitation, the blood lactate level of the rabbits in the resuscitation group was significantly decreased when compared with that of the control group, and the trend lasted until 90 min (p<0.01 for all). Moreover, at 30 and 60 min after the model was established, the lactate clearance rate of rabbits in the resuscitation group was significantly higher than that of the rabbits in the control group (p<0.01 for both; Table 4).

**DISCUSSION**

Traditionally, a large influx of liquid is allowed into the systemic circulation within a short period of time, and this increase in circulating blood volume is beneficial for the maintenance of arterial blood pressure. However, a sudden influx of liquid into the blood vessels will cause the infused blood to be retained in the venous system as the blood is in a state of stasis in the capillaries, which will cause a constant increase in intravascular hydrostatic pressure. In turn, this results in more intravascular liquid penetrating into the extravascular space through the blood vessel walls, which already show increased permeability due to the ischemic and hypoxic injury. Consequently, this process can cause edema of the internal organs, especially pulmonary edema which further aggravates the hypoxemia. In addition, this massive influx of liquid may also lead to heart failure, thus creating a vicious cycle. Further, after acute major bleeding, the peripheral veins will collapse, and phlebotomy or deep vein puncture will take a long time to complete. Hence, in some cases, the intravenous infusion rate cannot meet the needs of resuscitation. Under these circumstances, other procedures, besides intravenous infusion, are needed for rapid infusion, and arterial resuscitation provides a feasible option for emergency resuscitation.

There have been some reports on arterial blood transfusion. However, because of its complex nature and risk of adverse reactions such as vasospasm and limb ischemia, it is currently only used as a backup plan in cases of severe shock, acute blood loss, near-death state, and clinical death. In this study, during the process of preparing the hemorrhagic shock model, the average blood loss of the experimental animals was 25 mL/kg, and MAP decreased to 50% of the baseline value and stabilized at approximately 50 mmHg. MAP, pulse pressure, urine output, and blood lactate levels showed significant changes, indicating that the model met the requirements and was accurate. After the blood was partially reinfused by pulsed arterial reinfusion, MAP increased significantly, and in some rabbits, MAP could even be restored to the level before the shock. In addition, the pulse pressure also significantly increased and could be maintained for a long period of time.

It is generally considered that pulse pressure has a close relationship with tissue perfusion. In this study, the urine output in the resuscitation group was found to significantly increase as well, indicating that the renal perfusion of the experimental rabbits was partially restored. Furthermore, the blood lactate level of the experimental animals decreased, suggesting that hypoxemia, which appeared from the onset of the shock and was aggravated throughout the experiment, was restored to a certain degree.

Only after the oxygen debt is repaid, tissue acidosis can be corrected; for shock resuscitation to be considered complete, the aerobic metabolism needs to be recovered in addition to the hemodynamic parameters. Our results suggested that pulsed arterial blood reinfusion could increase pulse pressure, elevate tissue perfusion, and alleviate hypoxemia in a rabbit model of shock. Hypoxemia has always been considered the root cause of complications in shock and the major cause of mortality in these rabbits. Therefore, it is reasonable to believe that the application of pulsed arterial blood reinfusion may reduce early mortality in patients with hemorrhagic shock, while also having positive effects on the incidence of complications due to shock and late mortality.

In addition to rapidly increasing the effective circulating blood volume, arterial blood transfusion can pump the

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**Table 4.** Comparison of the dynamic changes in blood lactate levels and lactate clearance rates between the two groups treated with different reinfusion methods at different time points after establishing the rabbit model of hemorrhagic shock (Mean±SD)

<table>
<thead>
<tr>
<th>Group</th>
<th>Blood Lactate Levels (mmol/L [no. of animals])</th>
<th>Lactate clearance rate (% [no. of animals])</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>0 min</td>
<td>30 min</td>
</tr>
<tr>
<td>Resuscitation</td>
<td>2.85±0.15 (30)</td>
<td>5.16±0.42 (30)</td>
</tr>
<tr>
<td>Control group</td>
<td>2.96±0.47 (30)</td>
<td>5.13±0.51 (30)</td>
</tr>
<tr>
<td>t value</td>
<td>-0.661</td>
<td>-0.421</td>
</tr>
<tr>
<td>p value</td>
<td>0.517</td>
<td>0.678</td>
</tr>
</tbody>
</table>

The resuscitation group received pulsed carotid artery reinfusion, and the control group received pulsed intravenous reinfusion. Resuscitation began at 60 min. The value at 60 min was compared with that at 0 min (p<0.05). The number of animals is recorded within parentheses.
functions that were initially interfered with. As the blood and subcortical regions and thereby restoring the regulatory functions of the central nervous system, especially that of the cortex, this also alleviates the myocardial ischemia and facilitates the recovery of cardiac function. The blood that flows through the aorta quickly enters the kidneys, gastrointestinal tract, and other organs, thus increasing the blood perfusion, protecting the function of the internal organs, and slowing down the systemic inflammatory response syndrome, as well as reducing the risks of intestinal bacterial translocation and intestinal endotoxemia. Because the transfused blood in the artery flows retrogradely, it forms turbulences in the large vessels (similar to ventricle turbulence), which helps to mix the blood thoroughly. Therefore, the oxygen content of the blood transported to the various organs will not increase too abruptly, thus creating a buffer allowing for a gradual increase of oxygen content in the body. This helps reduce ischemia and reperfusion injury and can hence help reduce the risks of visceral and pulmonary edemas.

Moreover, the pressurized transfusion of arterial blood constantly stimulates the vessel walls, including the chemical receptors and baroreceptors of the aortic arch and carotid sinus. The reflex protection system of the body and the vagal reflex from the direct stimulation of blood on the arterial wall can result in a significant elevation of arterial pressure and lead to high blood pressure at the arterial end of the capillaries, which helps reduce capillary congestion. The decrease in heart rate induced by this reflex may increase the pulse pressure, improve the cardiac ejection capability, and eventually augment the tissue perfusion pressure; visceral vasodilation induced by this reflex also helps improve the visceral blood perfusion. A small proportion of the retrograde blood flows into the heart ventricle, thereby prolonging ventricular isovolumic relaxation and promoting rapid closure of the semilunar valves. As a result, the ventricular pressure will decrease sharply, thus forming a large suction force as the main driving force of rapid ventricular filling. This force will cause increased blood flow from the left atrium into the left ventricle, which helps alleviate venous congestion, and will also increase the left ventricular end-diastolic volume. Myocardial contractility is strengthened through self-regulation, and the diastolic extension is also conducive to coronary blood supply, thus forming a virtuous circle. As a result of pulsed arterial blood transfusion, the negative effect of stimulating the blood vessel wall is reduced and the occurrence of vasospasm is decreased. Mild protective vasospasm helps increase peripheral resistance to increase blood pressure and will not affect the ability of blood vessels to transport blood. Moreover, pulsed arterial blood transfusion may also help reduce endothelial injury, maintain endothelial secretion and barrier functions, reduce blood vessel wall inflammation, and reduce the risk of thrombosis. The pulsed procedure stimulates the aortic arch and chemical receptors to preserve them in a sensitive state, and therefore, resetting will not occur. Meanwhile, the pulsed procedure does not result in constant stimulation of the vagus nerve to cause hyperexcitability and will hence not cause a reduction in the cardiac ejection blood volume to aggravate low blood pressure. This reduces the blood pressure fluctuations as well as the interference on the hemodynamics and on the body’s own regulation during the buffering period.

Lastly, blood (especially autologous blood, including uncontaminated body cavity blood and frozen plasma extracted and prepared in advance from high-risk groups such as soldiers) has always been considered the best and most effective resuscitation solution. Apart from momentarily increasing blood volume and reducing apoptosis, the blood can moreover supplement coagulation factors, which help control bleeding.

Conclusion

With the recent advancements in arterial puncture technology, arterial resuscitation can enable direct transportation of blood to the heart, brain, kidneys, and other organs, thereby greatly reducing the reliance on cardiac function. Therefore, pulsed arterial resuscitation might be useful for the emergency treatment of hemorrhagic shock. However, further studies and discussions are still needed on whether pulsed arterial resuscitation can improve the clinical cure rate and reduce mortality.

Acknowledgments

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Conflict of interest: None declared.

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Tavşan hemorajik şok modelinde atımlı arteriyel resüsitasyon uygulaması

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²Hangzhou İkinci Hastanesi, Hangzhou Normal Üniversitesi Tıp Fakültesi, Nöroşirüj Bölümü, Hangzhou-Çin Halk Cumhuriyeti


BULGULAR: Kontrol grubuyla karşılaştırıldığında, 60. dakikada resüsitasyon grubunda OAB, nabız basıncı ve idrar çıkarımı anlamlı derecede daha yüksek (OAB: 83.67±3.90’a karşın, 38.19±3.50 mmHg, p<0.001; nabız sayısıındaki farklılık: 16.46±2.21’e karşın, 10.27±2.99 mmHg, p<0.001; idrar miktarı: 3.68±0.74’e karşın, 0.10±0.05 mL·kg⁻¹·dk⁻¹, p<0.001), serum laktat düzeyi ise anlamlı derecede daha düşüktü (3.82±0.50’e karşın, 6.49±0.61 mmol/L, p<0.001). Ayrıca resüsitasyon grubunda laktat kliрен orani anlamlı derecede daha yüksekti (30 dk: %0.26±0.11’e karşın, %0.25±0.14, p<0.001; 60 dk: %0.30±0.09’a karşın, %0.67±0.26, p<0.001).

TARTIŞMA: Hemorajik şokun acilen tedavisinde atımlı arteriyel resüsitasyon kullanabilir.

Anahtar sözcükler: Arter; hemorajik şok; nabız; resüsitasyon.