Assessment of resin perfusion in hepatic failure in vitro and in vivo

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Abstract

AIM: To observe the adsorbent effect of resin on endotoxin, cytokine, bilirubin in plasma of patients with hepatic failure and to determine the resin perfusion as an artificial liver support system in the treatment of hepatic failure.

METHODS: One thousand milliliters of discarded plasma was collected from each of 6 severe hepatitis patients treated with plasma exchange. The plasma was passed through a resin perfusion equipment for 1-2 h via extracorporeal circulation, and then absorbent indicators of transaminase, bilirubin, blood ammonia, endotoxin and cytokines were examined. In the meantime, study of in vivo resin plasma perfusion was performed on 7 severe hepatitis patients to compare the changes of endotoxin and cytokines in blood before and after perfusion.

RESULTS: The levels of total bilirubin, endotoxin, interleukin 1β and TNF-α in plasma were significantly decreased after in vitro resin plasma perfusion. The levels of interleukin 1β, TNF-α and endotoxin in blood were also evidently declined after in vivo resin plasma perfusion. Nevertheless, no obvious changes in IL-6, creatinine (Cr) and urea nitrogen (UN), blood ammonia and electrolytes were found both in vitro and in vivo.

CONCLUSION: Bilirubin, endotoxin and cytokines in plasma of patients with hepatic failure can be effectively adsorbed by resin in vitro. Most cytokines and endotoxin in plasma can also be effectively removed by resin in vivo. It demonstrates that resin perfusion may have good treatment efficacy on hepatic failure and can be expected to slow down the progression of hepatic failure.

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INTRODUCTION

In the research field of artificial liver, blood perfusion, blood filtration, plasma exchange techniques and methods are called non-biological artificial liver[1-3]. They are based on the mechanical mechanism of blood purification to remove toxins in body to achieve the treatment goal for hepatic failure[4-11]. New-style resin blood perfusion is mostly used in toxicosis rescue and uremia, etc in clinic, but was rarely reported in the treatment of hepatic failure. Since endotoxin and cytokine play a pivotal role in the pathogenesis of hepatic failure in severe hepatitis[12-16], it has become an important issue whether resin blood perfusion can effectively remove endotoxin and cytokine in body. Therefore, we examined the effect of a non-bioartificial liver with resin plasma perfusion on the plasma endotoxin and cytokine removal in patients with severe hepatitis to determine and assess the curative effect of resin plasma perfusion in the treatment of severe hepatitis.

MATERIALS AND METHODS

Subjects

Thirteen subjects with severe viral hepatitis (female 1, male 12, aged 28 to 57 years with a mean of 41.3 years) were enrolled in this study. All subjects belonged to chronic severe hepatitis caused by HBV infection according to the diagnostic criteria described in the Viral Hepatitis Protection and Cure Guideline established on a national conference. Of these patients, 8 were in middle stage, 5 in final stage, complication of encephalopathy, spontaneous peritonitis and hepatorenal syndrome was found in 11, 5 and 3, respectively.

Experimental protocol in vitro

One thousand milliliters of discarded plasma was collected from each of 6 severe hepatitis patients treated with plasma exchange, and mimic blood perfusion was performed in 3 h. The resin perfusion device (Lizhu Bio-material Company, Zuhai, China) and tubes were filled with 500 mL of 50 g/L glucose injection, then rinsed with 3 000 mL of normal saline containing 80 mg heparin from bottom to top at 50-100 mL/min. In the mean time, perfusion device was tapped by hand and rotated to exclude bubbles and particles. Plasma perfusion was performed for 1-2 h (average 1.7 h) and the velocity of plasma cycling through the perfusion device was at 80-120 ml/min. Plasma before and after circulation was collected, sealed and stored at -40 ºC. All the procedures, such as plasma collection, perfusion in vitro and sample harvesting used aseptic technique, syringe, cycling tubes and frozen tubes were irradiated by 60Co to remove pyrogen.

Experimental protocol in vivo

Resin plasma perfusion was performed in 7 patients with severe hepatitis. Temporal artery-vein circuit was established by
insertion of femoral and pectoral venous canulas. The patients were heparinized based on individual conditions. The first dose was 1-1.5 mg/kg, then 0.1 mg/kg each 30 min was added. The patients’ blood was introduced to a plasma separator (Fresenius Medical Care, Bad Homblurg, GER) at 100-120 mL/min for separation of plasma from the whole blood. The separated plasma was absorbed by the resin perfusion device, then mixed with separated blood cells, and returned to patients. The perfusion was maintained for 1-2 h (average 1.6 h) (Figure 2). The plasma before and after perfusion was collected and stored as the above methods.

Figure 2 Circuit diagrams of plasma perfusion.

Biochemical estimations and cytokines detection
Plasma aminotransferase, total bilirubin (TB), total bile acid (TBA), electrolyte, UN and Cr were measured using a biochemical autoanalyzer (Hitachi Co., Tokyo, Japan). Blood ammonia was measured by a blood ammonia detector (Shiga Co., Tokyo, Japan). Endotoxin was measured by using a limulus test kit according to manufacturer’s instructions (Shengxong Medical Technology Co., Shanghai, China). The level of IL-1β and TNF-α was measured by using a radio immunoassay (RIA) kit according to manufacturer’s instructions (Department of Radio Immunoassay, PLA General Hospital Technical Development Center ). IL-6 level was measured by using an ELISA kit according to manufacturer’s instructions (Shengxong Medical Technology Co., Shanghai, China).

Statistical analysis
All results were expressed as mean±SD. Comparisons between the groups and the same group before and after perfusion were analyzed by Student’ s t test.

RESULTS
Changes in transaminase and bilirubin
After resin perfusion device was used to absorb plasma of the patients with severe hepatitis in vitro, transaminase, TB and TBA were all declined with a statistically significant difference in TB, DB and IB before and after perfusion (t value was 2.081, 2.048, and 2.086 respectively, P<0.05).

| Table 1 Comparison of transaminase, TB and TBA before and after in vitro resin perfusion (mean±SD) |
|-------------------------------------------------|
| Before absorption | After absorption |
| ALT(IU/ L) | 74.17±9.68 | 47.83±22.33 |
| AST(IU/ L) | 88.63±5.58 | 58.33±4.64 |
| TB(µmol/ L) | 518.8±180.27 | 356.13±162.87 |
| TBA(µmol/ L) | 157.17±53.53 | 131.67±54.95 |

TBA was markedly declined in patients with in vivo resin plasma perfusion with a significant difference (P<0.01). Nevertheless, transaminase and TB had no marked change before and after absorption (Table 2).

| Table 2 Comparison of transaminase, TB, and TBA before and after in vivo resin perfusion (mean±SD) |
|-------------------------------------------------|
| Before absorption | After absorption |
| ALT(IU/ L) | 270.0±355.54 | 288.3±314.58 |
| AST(IU/ L) | 252.5±591.51 | 224.8±356.79 |
| TB(µmol/ L) | 519.66±209.00 | 460.8±165.99 |
| TBA(µmol/ L) | 194.8±47.32 | 166.7±48.98 |

Comparison of electrolytes and renal function
The electrolytes – K, Na, Cl, Ca, Mg, and P after absorption had no significant changes compared to those before absorption. UN and Cr were declined after absorption with no statistical significance compared to those before absorption with UN value of 6.79±4.14 mmol/L before absorption and 6.42±3.39 mmol/L after absorption, and Cr value of 135.67±30.77 µmol/L before absorption and 121.85±20.40 µmol/L after absorption.

Blood ammonia
After in vitro resin absorption, blood ammonia in plasma was not markedly decreased (98.17±20.60 µmol/L and 91.50±18.84 µmol/L). Similar to the result of in vitro, blood ammonia had no significant difference before and after in vivo resin plasma perfusion (48.25±8.63 µmol/L and 46.75±9.44 µmol/L).

Level of LPS and cytokines
The level of LPS was markedly decreased after in vitro absorption (t=6.604, P<0.01). The changes in TNF-α and IL-1β had a statistical significance (t was 2.876 and 3.673 respectively, P<0.05). IL-6 had no significant change before and after in vitro resin absorption (Table 3).

| Table 3 Level of LPS and cytokines before and after in vitro resin absorption (mean±SD) |
|-------------------------------------------------|
| Before absorption | After absorption |
| LPS (ng/ L) | 60.35±8.58 | 32.75±10.14 |
| TNF-α(ng/ L) | 1.49±1.06 | 1.04±0.91 |
| IL-1β(ng/ L) | 2.61±1.42 | 1.68±0.92 |
| IL-6(ng/ L) | 100.07±10.99 | 87.6±12.27 |

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| Table 4 Level of LPS and cytokines before and after in vivo resin absorption (mean±SD) |
|-------------------------------------------------|
| Before absorption | After absorption |
| LPS(ng/ L) | 58.64±11.03 | 36.13±5.24 |
| TNF-α(ng/ L) | 2.28±0.89 | 1.69±0.76 |
| IL-1β(ng/ L) | 1.98±0.82 | 1.24±0.88 |
| IL-6(ng/ L) | 88.41±24.96 | 84.85±16.55 |

\*P<0.01 vs before absorption.
DISCUSSION

Blood perfusion is a commonly used method for blood purification[17-20]. Activated charcoal representing perfusion equipment is most widely applied in the clinical treatment for drug toxicosis, uremia and liver coma and a good result has been obtained[21,22]. However, the shape of activated charcoal particles is irregular and their mechanical strength is inferior, thus particles are easily broken. If a broken particle directly contacts with blood, it will cause hemolysis and microvesSEL occlusion. Therefore, activated charcoal must be encapsulated before use. Synthetic resin is another kind of medical absorbent material with macropore high molecular polymers belonging to neutral macropore resin that absorbs substances via Vandar Val gravitation. Its absorbent ability has been found to be characterized by a fast absorbent speed, high mechanical strength, relative absorbent specificity mainly absorbing the substances with a molecular weight of 500-5000 Da and showing an outstanding absorbent ability to those toxins which can bind to proteins closely or are highly fat-soluble[23,24].

In the experiment of resin blood perfusion in vitro, we found that the resin blood perfusion device was able to effectively absorb transaminase and bilirubin but weakly affect kidney function and blood ammonia, considering that it absorbed medium molecules not small molecules. It illustrated the resin blood perfusion could improve the hepatic function in patients with severe hepatitis. In the meantime, it did not cause electrolyte imbalance and disturbance in the internal environment. Therefore, resin blood perfusion can be used as an artificial liver support system in the treatment of severe hepatitis.

Current studies considered that the pathogenesis of severe hepatitis was the superposition of virus-caused primary immunopathological damage and cytokine-induced secondary damage[25-27]. When liver barrier function was impaired, endotoxaemia would occur. Endotoxins stimulated the mononuclear phagocyte system inside and outside the liver, and thus, enormous cytokines were released. Furthermore, this cytokine-induced secondary hepatocellular damage played an important role in the course of hepatitis. Hereby, cytokine removal was good for alleviating liver damage, reducing leukoplasia and thrombocyte aggregates, maintaining internal environmental equilibrium, and sequentially slowing down or even reversing the progression of disease and improving prognosis[28-30].

Along with the development in molecular biology, research of antagonists for various cytokines has been launched, for example, monoclonal antibody, receptor antibody, soluble antibody. However, their clinical application is not ideal. In recent years, more and more researchers have begun to use blood purification to remove cytokines[31-33]. At present, it is still disputable whether blood purification can effectively remove cytokines[34,35]. Some researchers held that leucocytes could be activated by passing through the device during blood perfusion to cause the releasing of cytokines. In addition, it has been found the molecular weight of cytokines is relatively high, and cytokines are mostly bound to proteins in plasma, the half-life of cytokines is short and they are produced and metabolized quickly. Therefore, it would be hard to remove them via blood purification[36].

In this study, after plasma perfusion treatment for severe hepatitis, both TNF-α and IL-1β in plasma were significantly decreased, illustrating that plasma perfusion was an effective method for cytokine removal. The change of IL-6 was not significant, because IL-6 might be produced too fast, or it related to different absorbent efficacy of the perfusion device to various cytokines.

In conclusion, resin can effectively absorb bilirubin, LPS, TNF-α and IL-1β in vitro, and LPS, TNF-α and IL-1β can be significantly decreased in resin plasma perfusion in vivo, resin perfusion has good curative effects on severe hepatitis with hepatic failure.

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