Effects of a bioartificial liver support system on acetaminophen induced acute liver failure canines

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INTRODUCTION
Acute liver failure (ALF) is a well-known complication in clinical diseases and the current comprehensive treatment is not satisfactory. The recent developed artificial liver support systems, especially the bioartificial liver support systems (BALSS), which perfuse the ALF blood with the exotic hepatocytes to temporarily replace the complex functions of ill liver such as synthesis, detoxication and biotransformation, provide a brand-new approach to treat the ALF[1-8]. But the difficulties in acquiring high-density culturing of hepatocytes, lack of a stable and well-reproducible big animal model of ALF impeded the development of BALSS researches. In the present study, we have introduced the basic techniques of a hollow fiber bioartificial liver system, used the Chinese experimental miniature swines as the donor of hepatocytes, improved the culturing conditions of porcine hepatocytes in the bioartificial liver system and developed an acetaminophen-induced ALF canine model so as to assess the efficacy, safety and feasibility of the BALSS in treating the ALF canines.

MATERIALS AND METHODS
Animal
The healthy adult hybrid canines weighing 15kg and adult Chinese experimental miniature swines weighing 10kg, the male or female, were provided by the Experimental Animal Center of PLA General Hospital.

Reagent
All chemicals were obtained from Hong Kong TECA LTD Co. or Sigma Chemical Co.

The development of a canine model of ALF
The canines received multi-subcutaneous injections (sc) with acetaminophen (115mg/kg) to induce the ALF models.

The isolation and culturing of porcine hepatocytes
Hepatocytes were isolated from Chinese experimental miniature swines by collagenase digestion. Viability of the cells was assayed by trypan blue exclusion and AO/PI fluorescence staining.
The procedure of BALS treatment
Freshly isolated hepatocytes were cultured in the TECA-I BALSS from Hong Kong TECA LTD Co. and used to treat the ALF canines. The porcine hepatocytes were circulating through the exterior space of capillary hollow fibers in the BALSS. A circuit was developed between the femoral artery and vein on one side of the ALF canine and connected to the hollow fibers of BALSS. The canine’s blood circulated through the inner space of capillary hollow fibers and the substances exchanged through the membrane of hollow fibers with the porcine hepatocytes to achieve the perfusing function.

Experimental groups
Nineteen ALF canines were randomly divided into three groups: ① BALSS perfusion group (n = 9); perfused with the hollow fiber tube BALSS; ② drug group (n = 5): intravenous injection with arginine, sodium glutamate and acidi amino chaini branchi; ③ control group (n = 5): intravenous injection of 5% dextrose solution. Each treatment lasted six hours.

Biochemical assays
The plasma levels of ammonia, GPT, GOT, AKP and BUN were measured before and 2, 4, 6 hours, 1, 3, 5, 7, 14, 30 days after treatment.

Morphological observation
The liver, kidneys, heart, lungs were taken out when the canines died or survived for 30 days for paraffin section and HE stained and examined under light microscope.

Statistical analyses
The values of blood biochemical parameters were expressed as means±SD. Data between the three groups and pre-and post-treatment were compared using t test. Statistical significance was considered when P<0.05.

RESULTS
The development of a canine model for acetaminophen-induced ALF
After injected with acetaminophen, the thirty-eight healthy canines (21 male, 17 female) demonstrated a significantly increased level of NH₃, GPT, GOT and AKP at 48 hours (P<0.01, Figure 1) while plasma concentration of BUN had no change. In the mean time, symptoms appeared such as vomiting, food refusal and listlessness. Histologic examination showed massive necrosis of the hepatic tissues. Sixty-three per cent had ALF, in which the plasma levels of NH₃ and GPT were higher than 100 µmol/L and 60U/L respectively.

The porcine hepatocytes
Using the modified enzymatic digestive method, each liver yielded approximately 0.8 - 5×10¹⁰ hepatocytes. Viability of the final suspensions averaged 60% - 80% by trypan blue exclusion and AO/PI fluorescence staining.

The liver function recovery of ALF canines after treatment with BALSS
All ALF canines in the control group, with increasing levels of plasma ammonia, GPT, GOP and AKP died 2 - 30 hours after treatment. All the five ALF canines in drug group died 6 - 36 hours after treatment. The plasma levels of ammonia, GPT, GOT and AKP declined slightly during the treatment, but it had no significant difference compared with that in the control group (P>0.05). In BALSS perfusion group, all nine ALF canines showed remarkable liver function recovery. During BALSS treatment, the levels of plasma ammonia, GPT, GOT and AKP began to decline at the 2nd hour and the declination was even more obvious compared with pre-treatment at the 4th hour and 6th hour (P < 0.01) (Figures 2, 3, 4, 5). The levels of these biochemical parameters had significant differences at the 6th hour of treatment between BALSS perfusion group and the drug group (P<0.01). Except three canines in BALSS perfusion group who died 6 and 12 hours and 5 days after treatment because of narcotization and wound bleeding, the other 6 canines survived more than 30 days.

The morphological changes of liver in ALF canines after treatment with BALSS
The liver, heart, kidneys and lungs were taken out for pathological examination when the canines died and survived for 30 days after treatment. The liver tissues of the dead ALF canines showed massive necrosis in the drug and control groups. Thirty days after treatment, the liver tissue of ALF canines in BALSS perfusion group had no obvious hepatocyte necrosis but a slight cholestasis and fibrosis.

![Figure 1](image1.png) The changes of canines’ biochemistry parameters before and after injections of acetaminophen.
ALF is usually an integrative response caused by viruses, drugs and toxins which leads to severe hepatocyte injury and the rapid loss of synthetic, metabolic and regulative functions, followed by the release of non-inflammatory cellular factors and other indefinite toxins into the circulation. Its mortality was as high as 50% - 80% and no effective clinical therapy is available currently. Because the iled liver has a tendency of reversibility, a reproduction can occur if we replace the hepatic functions by artificial methods. Since the liver has more than 500 complex physiological functions, the current non-biological ALS such as activated charcoal absorption can not replace the liver functions completely although it can clear some middle and small molecular weight substances in the circulation. In BALSS the exotic hepatocytes were used to exchange the substances with the ALF blood, so that it can temporarily replace the complex liver functions of ALF patients directly from the exotic hepatocytes. Now there are several types of BALSS and a few small animal experiments and clinical study reports with various therapeutic effects[1-7,9-11]. The studies of BALSS in China are still in the initial stage and the BALSS techniques are not mature. This experiment studied the main problems such as the source and culturing of hepatocytes, the development of ALF canines model and a ssessment of the efficacy and safety of BALSS treatment based on the advanced techniques of hollow fiber tube BALSS introduced from abroad.

A large quantity of hepatocytes with a well-functioned condition and high-density in culture in the BALSS is a key factor in success. Some authors used the human liver tumor cell lines, human embryo hepatocytes and porcine hepatocytes as the sources of hepatocytes[9-11]. But the possible dangers of the tumor cells to the receptors and the scarce of the embryonic cell hampered the studies of BALSS. Our experiment selected the quarantined pure strain Chinese experimental miniature swine as the source of hepatocytes. Using the modified enzymatic digesting and culturing method, we acquired a large quantity of porcine hepatocytes that can meet the needs of BALSS. The development of a large animal ALF model, especially with high-reproductivity, stablity and a similar pathological changes to ALF patients, is still a major difficulty. Some people established the ALF canine models by surgical operation to occlude the blood supply for liver temporarily[12]. But this kind of models can not suit the studies of ALF or drugs because of the multiple organs failure induced by ischemic-reperfusion injury and the variability of death time. We used the multiple injections of acetaminophen to
induce the ALF canines. Forty-eight hours after injection, the blood hepatic biochemistry parameters had obvious changes. The canines’ liver tissues manifested massive necrosis. The ALF canines would die within 56 - 84 hours without treatment. This disease model had a good reproducibility and the changes of blood biochemical parameters were similar to ALF patients except the plasma BUN. The death time of ALF canines is relatively stable and suits the studies of BALSS and drugs.

In our experiment, we developed an ALF canine model induced by acetaminophen and then treated with the clinical routine drugs, BALSS and infusion of dextrose solution for evaluating the efficacy, safety and feasibility of BALSS to treat the ALF canines. The biochemical parameters related to hepatic functions of the ALF canines in the control group were elevated except for the plasma BUN. The canines died within 2 days. The drug therapy could partly correct the changes of biochemical parameters of the ALF canines, but with a very limited efficacy, and the canines died 2 days after treatment. The perfusion treatment with the BALSS of cultured Chinese experimental miniature porcine hepatocytes lasted 6 hours. The blood biochemical parameters of hepatic functions of the ALF canines were corrected obviously at the 4th hour of perfusion. The effect was significantly superior to which used in the other two groups. ALF canines can survive more than 30 days after perfusion therapy. The pathological changes of liver were also recovered obviously. The results showed that the porcine hepatocytes in this kind of BALSS could temporarily replace the hepatic functions of ALF canines by perfusion through the membrane of the hollow fiber tube, and then offered a chance for regeneration and repairment of the canine liver. It is suggested that the BALSS could temporarily replace the hepatic function of ALF patients safely and effectively, and may become a brand-new method for the treatment of ALF.

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