An Experimental, Non-Uremic Rabbit Model of Peritoneal Dialysis

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Summary
Peritoneal dialysis (PD) is a well established method of depuration in uremic patients. Standard dialysis solutions currently in use are not biocompatible with the peritoneal membrane. Studying effects of dialysate on peritoneal membrane in humans is still a challenge. There is no consensus on the ideal experimental model so far. We, therefore, wanted to develop a new experimental non-uremic rabbit model of peritoneal dialysis, which would be practical, easy to conduct, not too costly, and convenient to investigate the long-term effect of dialysis fluids. The study was done on 17 healthy Chinchilla male and female rabbits, anesthetized with Thiopental in a dose of 0.5 mg/kg body mass. A catheter, specially made from Tro-soluset (Troge Medical GMBH, Hamburg, Germany) infusion system, was then surgically inserted and tunnelled from animals' abdomen to their neck. The planned experimental procedure was 4 weeks of peritoneal dialysate instillation. The presented non-uremic rabbit model of peritoneal dialysis is relatively inexpensive, does not require sophisticated technology and was well tolerated by the animals. Complications such as peritonitis, dialysis fluid leakage, constipation and catheter obstruction were negligible. This model is reproducible and can be used to analyze the effects of different dialysis solutions on the rabbit peritoneal membrane.

Key words
Peritoneal dialysis • Experimental model • Non-uremic rabbit

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Introduction
Peritoneal dialysis (PD) is a well-established method of depuration in the end-stage renal disease patients. The PD system consists of three main components: peritoneal microcirculation, peritoneal membrane and the dialysate compartment. The peritoneal filtration membrane is composed of peritoneal mesothelial cells, mesothelial basement membrane, connective tissue, blood vessels basement membrane and endothelium. This living structure is more a functional barrier than strictly an anatomical formation.

Standard dialysis solutions currently in use are not biocompatible with the peritoneal membrane. They contain glucose in high, non-physiological concentrations, acting as an osmotic substance. Furthermore, the dialysate contains lactates which maintain its low pH, thus contributing to its negative effects on the peritoneal tissue. During sterilization and preservation of the solute, glucose degradation products and advanced glucose degradation products are formed, which adversely affect the peritoneal structures. The high glucose concentration causes non-enzymatic glycosylation of tissue proteins. This may explain the resemblance of morphological alterations in peritoneal microcirculation during long-term dialysis to those found in blood vessels of diabetic patients: loss of the mesothelial layer, thickening of submesothelium due to increased deposition of collagen and hyaluron in interstitium, interstitial fibrosis, thickening of mesothelial basement membrane and endothelial basement membrane.
of small peritoneal blood vessels accompanied by neoangiogenesis (Stojimirovic et al. 2001). Histological changes correlate with dialysis duration and frequency of the use of dialysate with high glucose concentration. Structural changes affect the quality of dialysis, as they increase the velocity of low molecular mass solute transport, increase peritoneal microvascular surface and decrease ultrafiltration rates.

Studying the effects of dialysate on peritoneal membrane in humans is still a challenge due to ethical and technical limitations, as tissue samples can be obtained solely when placing or removing the peritoneal catheter. Therefore, various experimental models have been developed in order to investigate the impact of dialysis solutions on peritoneal tissue (Ter Wee et al. 2003). There is, however, still no consensus on the ideal experimental model (Topley 2005, Mortier et al. 2005).

We therefore aimed to develop a new, experimental, non-uremic rabbit model of peritoneal dialysis, which would be practical, easy to perform, inexpensive, and convenient to study the long-term effect of dialysis fluids.

Methods

Animals

The study was performed on 17 healthy Chinchilla male and female rabbits, weighing 2739±388 g at the beginning of the experiment.

The rabbits were kept in individual cages and received standard rabbit pellets (Veterinary Institute, Subotica, Serbia) and water ad libitum. All rabbits were allowed to adapt to the new living conditions for at least five days prior to catheter insertion. Eleven rabbits (6 male and 5 female) completed the entire planned experimental procedure, i.e. 4 weeks of peritoneal dialysate instillation. During the study period of five weeks (one week for recovery following catheter placement, and four weeks of dialysis) a diary of animal behavior was kept. It included the following data: body mass, body temperature, food intake, diuresis, defecation, antibiotics administration, other therapy and interventions if necessary (wound toilette, catheter suturing etc.).

Sample collecting

Peritoneal tissue is extremely fragile and susceptible to mechanical irritation and environmental factors. Collecting tissue samples was therefore performed strictly respecting the guidelines from literature (Di Paolo et al. 1996, Gotloib and Shoshtak 2000, Williams et al. 2002). Ellipsoid tissue samples, 18 mm x 3 mm, of parietal peritoneum were taken immediately after opening the abdomen (Dobbie 1993). The tissue was fixed straight away for 24 h in 10% formaldehyde with 0.1 M Sorensen’s phosphate buffer pH 7.4, dehydrated in 96% ethanol, then routinely processed for embedding in paraplast, and stained with hematoxylin-eosin and toluidin-blue to be analyzed by light microscopy (Opton Photomikroskop III).

For transmission electron microscopy (TEM) tissue samples were fixed for 24 h in 4% glutaraldehyde with 1% tanine acid to ensure better fixation of membrane structures. Without tanine acid fixation lasted from 24 hs to several days. Fixatives were diluted in cacodylate or Sorensen’s phosphate buffer 0.1 M, pH 7.4. The tissue was then rinsed three times for 10 min with the same buffer, and then three times for 10 min in cacodylate buffer. Samples were postfixed with 1% osmium tetroxide in 0.1 M cacodylate buffer, pH 7.4 and left overnight in 4% uranyl acetate. They were then dehydrated in ethanol and propylene-oxide and embedded in Epon. Fine sections were contrasted with uranyl acetate and lead-citrate (Hayat 1986) and analyzed with a transmission electron microscope (Philips M208S).

Anesthesia procedure

Animals were anesthetized according to an existing protocol which was adapted during the study. For catheter implantation and removal rabbits were anesthetized intravenously with Thiopental BP 1G (Rotexmedica, Trittau, Germany) 1 ng/kg body mass via an ear vein. After losing two animals with this protocol, other rabbits were anesthetized with the same anesthetic (Thiopental injection BP 1G), via the same route, but the dose was halved to 0.5 mg/kg body mass. This dose of anesthetic was sufficient for adequate catheter implantation or removal and showed no adverse effects on the animals.

Surgical procedure

Catheter placement was a modified version of the procedure described by Zweers et al. (1999). At the beginning of our study we used double-lumen central venous catheter (Arrow International Inc. USA Product No. CV-17702-E) as a peritoneal catheter. Implantation of this catheter was performed as described in the following text.
When anesthetized, the animals were shaved on the back of the neck between the ears and scapulae, as well as below the left costal arch (upper left hemiabdomen). The surgical field was prepared in a standard way. Then, a longitudinal incision, 3 to 4 cm long, was made with a scalpel, starting 2-3 cm distally from the left costal arch edge, 4-5 cm from median abdominal line and in parallel to it.

After the skin was cut, subcutaneous space was entered either sharply or partly bluntly, and a tunnel was made from the abdomen to the neck area, with a thoracic drain No. 16 mandren (Figure 1a). The catheter was made to exit in the dorsal part of the neck, between the ears. The double-lumen central venous catheter was then pulled over the mandren and carefully pulled back, through the tunnel, into the abdominal region. Abdominal muscles were then sharply or bluntly moved apart to access the peritoneum. Immediately after opening the peritoneal cavity, biopsies of peritoneal tissue were taken from diagonal edges and the catheter was placed on the floor of the peritoneal cavity.

Peritoneum was sutured with a continuous suture using Vicril 4-0. Part of the catheter was fixed to it. Muscles were sutured with chromium Cutgut 3-0, and fascias with ongoing suture using Vicril (Dexon) 3-0. The skin was closed with single sutures. The catheter was fixed in the tissue at its entrance and exit sites. A sterile gauze was placed at both sites and fixed with a bandage wrapped around the body.

The double-lumen central venous catheter (used as the peritoneal catheter) was placed in the described manner in four animals. In the first few postoperative days, however, the rabbits bit off the external catheter branches and even pulled out the whole catheters, even though they were fixed at both ends. Due to their length, branches of the catheter stretched over animals' ears, irritated them and were easily accessible. These catheters could not be shortened as they had an original extension for attaching the infusion systems on their exit end.

This procedure, therefore, had to be modified. During the further course of our study we used a part of the infusion system Tro-soluset (Troge Medical GMBH, Hamburg, Germany), initially shortened to 45 cm, as a dialysis catheter. Anesthesia, shaving, operative field preparation, cutaneous incision and tunneling were all performed as previously described. The infusion catheter, 45 cm long, was pulled over a mandren (Fig. 1b) and led through the preformed tunnel (Fig. 1c). Although the catheter was made of very soft plastic, the intraperitoneal part was sharply cut and we protected the apical part of the catheter with 1 cm of soft rubber from an infusion set. The rubber was pulled over the catheter apex so that it did not make any contact with the peritoneal cavity and its organs. Then, four holes (2-3 mm in diameter each) were made with surgical scissors at the end part of the catheter, near the rubber cap (Fig. 1d). The peritoneum was then accessed as previously described. The catheter (shortened and adjusted to the animal's size) was placed on the floor.
of peritoneal cavity (Fig. 1e). The exit site of the catheter was at the back of the neck, between the ears (Fig. 1f). The wounds were then closed as previously described.

For catheter removal a new incision, 2-3 cm long, was made in front of the previous one. We opted for this approach because of the existing scar and to avoid possible damage to the peritoneum when collecting tissue samples. The same access to the peritoneum was made as the one previously described. Two tissue samples, one for light and one for electron microscopy, were taken. The catheter was then identified. In four animals it had first to be carefully freed from the omentum wrapped around its tip. The rubber apical part of the catheter was excised and the catheter was pulled out through the tunnel at its exit site on the dorsal part of the animal's neck. The abdominal wound was then closed as previously described.

**Infection prevention**

Infection was prevented with daily injections of cefuroxime (Nilacef®, Hemofarm and GlaxoSmithKline, England). Three days prior to catheter implantation and three days following catheter removal the antibiotic was given intramuscularly twice a day, in a daily dose of 150 mg. When peritoneal dialysate instillations were started, cefuroxime was administered intraperitoneally, through the catheter, once a day, in a daily dose of 75 mg. In the case that infection was suspected, gentamicin (Gentamicin®, Zdravlje, Leskovac, Serbia) was added intramuscularly, in a daily dose of 2 mg.

**Dialysis procedure**

We used dialysis solution (Dianeal PD4 Glucose, Baxter Vertriebs GmbH, Wien, Austria) with 3.86 % w/v glucose concentration preheated at 37 °C. Dialysis solution was first instilled on the seventh day following catheter placement. Instillations were then continued daily for the total of 28 days. The full instilled dose of dialysis solution was 40 ml/kg of body mass. In order to avoid respiratory problems caused by introduction of large amounts of fluid, observed at the beginning of instillation, animals were first administered 60 ml of the dialysate. This quantity was then increased by 10 ml each following day until it reached 40 ml/kg.

Daily heparinization was performed in order to prevent catheter clothing. After insertion, the catheter was heparinized with daily injections of 10 IU heparin sodium (Heparin®, Galenika a.d, Beograd, Serbia).

**Statistical analysis**

Statistical analysis of our data was made in Origin 7.0 and Microsoft Office Excel 2003. Results were presented as mean ± S.D.

**Results**

Eleven of the 17 rabbits enrolled in the study concluded the 5-week follow up, namely one week for recovery following the catheter placement and 4 weeks of daily dialysate instillation. Four rabbits were excluded from the study due to catheter loss (three of them had a double-lumen central venous catheter and one had a new, improvised catheter inserted). One rabbit died from anesthesia during peritoneal catheter implantation and one rabbit died one day following the catheter implantation. Thus 64.5 % of animals concluded the 5-week follow up, with a 89 % survival rate.

The reduced dose of anesthetic, as described previously, was sufficient for adequate catheter implantation or removal and showed no adverse effects on the animals. The rabbits made complete recovery immediately after surgery.

The instillation of a full dialysate dose at the beginning of the follow up resulted in a transient episode of dyspnea in the first rabbit treated. Dyspnea spontaneously resolved in a couple hours. This problem was exceeded by starting the instillation with 60 ml of dialysate on the first day of treatment, then increasing the amount of dialysate by 10 ml per day until reaching the full dose. The rabbits were instilled with full dialysate
dose over 5 to 7 minutes with excellent tolerance.

During the follow up, the rabbits had mostly normal body temperature (Fig. 2) and gained body weight (Fig. 3).

Only one episode of peritonitis was suspected in one rabbit which developed diarrhea on the fourth day of dialysate instillation. Although its body weight loss was only 100 g, i.e. less than 5% of total body weight, and fever was not higher than 39.5 °C, the animal was treated with cefuroxime intraperitoneally and gentamicin intramuscularly for 7 days. Diarrhea resolved after three days of treatment with the antibiotics.

All rabbits had daily urination and defecation. The only exception was a rabbit with constipation (starting from day 12 after peritoneal catheter placement). This animal received 20 ml of soap water (per rectum) and after this treatment it had daily defecation and a stable body weight.

In one rabbit we noticed subcutaneous dialysate leakage in the abdominal region on the twenty-seventh day of the follow up. The leak was reduced by slowing the rate of dialysate instillation.

**Discussion**

Both the uremic state and chronic exposure to PD fluid are associated with a development of functional and structural alterations of the peritoneal membrane (Stojimirovic et al. 2001, Trpinac et al. 2002). Commercial glucose-based peritoneal dialysis solutions expose peritoneum to hyperosmolar glucose, containing a variable amount of non-enzymatic glucose degradation products. Long-term peritoneal dialysis with glucose-based dialysis solutions has been associated with diabetes-like alterations of peritoneal tissue.

Animal models are important for understanding the physiology and pathophysiology of peritoneal structural and functional alterations during peritoneal dialysis. In vivo studies on animal models permit analysis of dialysis solutions biocompatibility under the conditions that allow interactions between dialysate and peritoneal membrane, as well as dynamic changes in dialysate composition that mimic the clinical situation closely (Zweers et al. 1999, Gaggiotti et al. 2005). Experiments on animal models showed similar morphological alterations of peritoneal membrane as in humans, but after a much shorter time of exposure to bioincompatible dialysis solutions (Topley and Lameire 2001). The toxic effect of bioincompatible dialysis solutions on all peritoneal cell types was proved in in vivo experiments. A large variety of animal models of PD might be a reason why there is still no consensus on the methodology and approach and no ideal model for studying PD.

Rats and rabbits are the most common animal models for PD. Rats are cheap and easy to breed.
limitations of the rat model are a consequence of rodent physiology. High serum amylase levels in rats are accompanied by high intraperitoneal amylase concentrations. Since amylase causes local degradation of icodextrin followed by a rise in dialysate osmolality, the rat model is inappropriate for evaluation of icodextrin effects (Mortier et al. 2005). Rabbits survive longer than rats on dialysis (Garosi and Di Paolo 2001). The ratio of peritoneal surface area and volume exchange in rabbits and humans is similar (Mortier et al. 2005). The disadvantage of rabbit model, however, is the fact that rabbits are very sensitive animals, difficult to keep.

In order to study PD induced alterations on peritoneal tissue we modified the rabbit model of peritoneal dialysis (Schambye et al. 1992). We have chosen this animal because it was more suitable for housing, feeding and keeping conditions, as well as the possibility of surgical treatment (surgical instruments, improvised peritoneal catheter, biopsies) available to us.

Different methods for fluid instillation during PD on experimental models are used by different authors. In some studies the test fluid was directly injected into the peritoneal cavity. In others, development of custom-made miniature peritoneal catheters allowed the so-called «open» PD system, with possible instillation and removal of the PD fluid (Lameire et al. 1998, Pawlaczyk et al. 2001). Another, the so-called «closed» system, uses a permanent catheter tunneled from the peritoneal cavity to the neck of the animal. In this case, drainage of the dialysate is not possible (Margetts et al. 2001, Zareie et al. 2001, Mortier et al. 2004).

Since we did not have an original peritoneal catheter for animals available, we opted for an improvised one. Using a part of the infusion system as a peritoneal catheter proved to be a fairly good option. Animals tolerated it well and it did not cause infection. Dialysate instillations were easily performed, and even dialysate removal was possible, which should enable studies of membrane function in the future. A special advantage of this model is that animals were unrestrained and awake, they had free access to food and water, they tolerated all procedures well and did not lose weight.

The most important technical problems in animal models of chronic PD are frequent obstruction of peritoneal access and development of infection. We would like to stress that during the five-week period of follow up no catheter obstruction occurred among 11 animals in our study. Use of heparin, besides desirable anticoagulant effects, is followed by undesirable effects such as modulation of inflammatory cell activity, proliferation of the cells, synthesis of extracellular matrix and neoangiogenesis (De Vriese et al. 2001). Although we recognize these side effects of heparin, we still prefer to use it rather than to perform omentectomy. As heparin is also used in clinical practice when problems with catheter functioning occur, its usage in animal models actually mimics real-life situations. The use of heparin-coated catheters, however, seems to be the method of choice of gaining peritoneal access (De Vriese et al. 2002).

Frequency of dialysate instillation varies among investigators from once to twice or even three times a day. The results suggest that severity of alterations correlates with the number of exchanges, regardless of the overall dialysis time.

Definition of peritonitis in animal models is still arbitrary. Most often used criteria for peritonitis diagnosis are positive dialysate culture and dialysate WBC count higher than 1000 cells/mm³ (Mortier et al. 2003, 2004). Peritonitis is suspected when there is a more than 5 % daily body weight loss, fever and diarrhea. Different strategies are in use to prevent and treat peritonitis (Zhou et al. 2005). Experimental data support prophylactic administration of antibiotics during the entire study period (Mortier et al. 2003). Prophylactic administration of cefuroxime successfully prevented intraperitoneal infection in animals used for our model of chronic peritoneal dialysate exposure. Only one rabbit showed clinical signs of peritoneal infection, which was successfully cured with an additional antibiotic.

Conclusions

The presented non-uremic rabbit model of peritoneal dialysis is relatively inexpensive, does not require sophisticated technology and was well tolerated by the animals. Complications such as peritonitis, dialysis fluid leak, constipation or catheter obstruction were negligible. This model is reproducible and can be used to analyze the effects of different dialysis solutions on rabbit peritoneal membrane.

Conflict of Interest

There is no conflict of interest.

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