A Novel three Cuff Peritoneal Dialysis Catheter with Low Entry Technique: Three Years Single Center Experience

Abstract

Objective: To share our 3-year experience with the new three-cuff peritoneal dialysis (PD) catheter with the low-entry technique and to study its effect on infectious and non-infectious complications as well as its impact on catheter survival.

Methods: This is an observational study which was carried out in a university hospital over 3 years. The study involved 153, three-cuff PD catheter insertions in 150 incident PD patients. The study was carried out in our PD center and extended from December, 2012 till January 2016 with a mean follow-up period of 15 months. All patients used automated peritoneal dialysis (APD). Throughout the study, we analyzed survival rate, functionality and complication profile of our new catheter.

Results: Four patients had inguinal hernia and 1 had omental wrapping. Catheter migration, however, was 0.0% with our 3-cuff PD catheter using our new technique. A total of 25 catheters had to be removed. Indications for catheter removal were successful transplantation (n = 7), hernia (n = 4), omental capture (n = 1), ultrafiltration failure (n = 2), psychological causes (n = 4), abdominal surgery (n = 1), severe tunnel infection (n = 3), and unresolved peritonitis (n = 3). The rate of peritonitis was as low as 0.106 per patient-year equivalent to 1 episode of peritonitis per 112 patient-months. At the end of the study, catheter survival was 91.3%.

Conclusion: The low entry-site of our PD catheter seems to prevent catheter migration. The 3-cuffs probably act as an additional safeguard against peritonitis.

Keywords: Peritoneal dialysis; Three-cuff PD catheter; Migration; Peritonitis; Catheter survival

Introduction

Peritoneal dialysis is an established form of renal replacement therapy for end stage renal disease. Catheter malfunction developed in 12.7%-35% of patients with Tenckhoff catheter. It is one of the main causes of peritoneal dialysis technique failure [1,2]. The most common causes of catheter malfunction are peritonitis, catheter tip migration and omental wrapping [3]. Indeed, both early and late catheter complications have limited catheter survival to only 51%-60% at 18 moth, 35%-51% at 24 months and 22%-33% at 36 moth [4,5]. Uncomplicated access to peritoneal cavity using permanent indwelling catheter is the key factor for successful peritoneal dialysis. Since the introduction of Tenckhoff catheter different catheters’ designs were introduced to prevent catheter displacement. Examples are the oreopoulos, the swan neck, the valli, the cruse, and the ash [6-12]. In 1992 Dipolo et al. [13] invented self-located catheter aiming at reducing catheter migration. They added material with high specific weight (tungsten cylinder at the distal end of the catheter) that tends to gravitate especially when the patient is erect. Although different studies showed low incidence of migration with the self-located catheter, others reported some drawbacks. Roberto Russo of the Bari group [14], reported increased incidence of adhesions (viscero-omental and viscero-visceral) that complicated removal of the self-located catheters.

This complication is most likely due to the sudden increase in diameter between the silastic and the tungsten cylinder. It could also be due to the presence of a foreign body in the peritoneal cavity causing a peritoneal reaction leading to the formation of inflammatory scar tissue. In addition, the cost considerations might be important; with costs amounting to approximately US$310, the self-locating catheter is about 70% more expensive than the simple straight Tenckhoff catheter [15]. There is no consensus in the literature concerning the superiority of one particular catheter design and length over the other and each method of catheter insertion has its benefits and proponents, but no technique has been shown to be preferable over the other [16]. We, at King Fahd Hospital of the University, developed a new PD catheter and a new technique for laparoscopic catheter insertion. The new catheter is three-cuffed and the new method totally eliminated the possibility of catheter-tip migration and significantly reduced the incidence of peritonitis in addition and hence improved catheter technical survival [17]. The objective of the current study was to share our 3-year experience with the 3-cuff low-entry-site new PD catheter that demonstrated zero migration, low peritonitis rates and high catheter survival.

Material and Methods

Using our new technique, a total number of 153, three-cuff PD catheters were placed in 150 incident PD patients at King Fahd
Hospital of the University, PD center. The study extended from December 2012 throughout January 2016. Exclusion criteria were pregnancy, morbid obesity, ongoing or previous peritonitis and patients’ refusal. Table 1 summarizes the characteristics of patients’ population included in the study. The study was conducted in accordance with the revised Declaration of Helsinki, and the study protocol was approved by the local Research Ethics Committee. All patients were above the age of 18 years and a written informed consent before participation in the study was required from every patient after full explanation of the new procedure. All catheters were evaluated for infectious and mechanical complications as well as technical survival.

Table 1: Demographic characteristics of APD patients.

|                        |        |
|------------------------|--------|
| Age, years (mean±SD)   | 52.2±18.7 |
| Gender (F/M), n (%)    | 80(53.3)/70(46.7) |
| DM, n (%)              | 61(40.7) |
| HTN, n(%)              | 88(58.7) |
| BMI, mean              | 26.8±2.5 |
| Total APD, mean (months) | 15.4±5.8 |
| GFR, ml/min (mean)     | 8.7±2.0 |
| Serum BUN, mmol/L (mean) | 28.4±11.5 |
| Serum Cr, µmol/L (mean) | 838.5±31.3 |
| Serum albumin, g/dl (mean) | 3.6±1.4 |
| Hgb, g/dl (mean)       | 9.1±1.8 |
| Primary renal disease, n(%) | 47 (31.3) |
| Diabetic nephropathy   | 28 (18.7) |
| Glomerulonephritis     | 15 (10.0) |
| Hypertensive renal disease | 8 (5.3) |
| Interstitial nephritis | 14 (9.3) |
| Systemic disease       | 14 (9.3) |
| Hereditary renal disease | 8 (5.3) |
| Unknown                | 14 (9.3) |

DM: Diabetes Mellitus; HTN: Hypertension; BMI: Body Mass Index; GFR: Glomerular Filtration Rate; BUN: Blood Urea Nitrogen; Cr: Creatinine; Hgb: Hemoglobin

Catheter Description

The Saudi PD catheter is silicone-rubber and about 57 cm in lengths. Details of the Saudi PD catheter are shown in Figures 1 and 2 A. Further details were demonstrated in a previous publication [16].

Surgical technique for the triple-cuff catheter insertion

The procedure was carried out with patient in the supine position under general anesthesia using aseptic precautions. A Veress needle was used to create pneumoperitoneum at pressure of 10-12 mmHg. A 5-mm port was inserted in right hypochondral region at middavicular line, 2cm below the costal margin for laparoscopic camera (30 degree). Diagnostic laparoscopy was then performed to rule out adhesions or herniations (Figure 2-B & 2 C). The operating table was then placed in about 30 degree trendelenburg position. Under direct vision, the three-cuff Saudi catheter was passed caudally through the pull-apart sheath over a 90-cm stylet into the peritoneal cavity. Above the symphysis pubis the tip of the catheter was placed in the true pelvis (towards the urinary bladder), with the distal cuff in the rectus sheath before removing the stylet. The distal cuff of the catheter was secured with purse-string suture on the fascia anterior to the rectus muscle by using 2/0 absorbable vicryl stitches. The tip of catheter was in the pouch of Douglas and the rectovesical pouch in female and male respectively. The pull-apart sheath was removed, leaving the catheter in the peritoneal cavity (Figure 2-B, 2-C and 2 D). Following that, a subcutaneous tunnel was created for the catheter with selection of a midway point at the umbilico-crestal line to be the output of the catheter. The end of the catheter attached to the stylet was advanced into the tunnel and pulled out from the above-mentioned point; the second cuff about 10 cm from the distal one and the proximal cuff 2 cm from the exit site. Figure 2-D and E illustrates the final position of the 3-cuff PD catheter. The function of the catheter was checked by flushing normal saline to rule out kinking or obstruction. The skin incisions of the camera port and entrance were sutured. Xylocaine with adrenaline diluted in normal saline was injected in the incision site and in the tunnel space.
Antibiotic prophylaxis

Was given with first-generation cephalosporin antibiotics administered intravenously prior to the procedure. Automated peritoneal dialysis (APD) was generally instituted 14 days after PD catheter insertion. Patient training was performed during this period with low volume exchanges.

Exit-site care

Routine exit-site care by the patient consisted of daily washing with anti-bacterial soap, thorough drying, cleaning with iodine solution and application of a small amount of mupirocin ointment (approximately 10 mg, a 1/4 inch dab) around the catheter exit-site using a cotton swab. Catheters were anchored with tape and small gauze dressing to prevent exit-site trauma. The exit-site was examined by the physician and a trained nurse at each clinic visit. Signs and symptoms of exit-site infection (ESI) and/or peritonitis were explained to the patients at each visit, and they were instructed to report to the unit once infection was suspected or occurred. Swabs and cultures were obtained from the drainage, from around the catheter and from any exit-site discharge. The diagnosis of PD peritonitis was made by the presence of cloudy effluent dialysate, abdominal pain with or without fever, and more than 100/mm3 of white blood cells with over 50% polymorphonuclear leukocytes in peritoneal effluent over 2 h.

Statistical analysis

The statistical analysis was performed using the IBM SPSS statistics Version 20 (IBM, Inc., New York, USA). Continuous variables were presented as means±SD and qualitative variables were presented as frequency and percentage. Catheter survival was analyzed by the Kaplan-Meier method.

Results

From December 2012 throughout January 2016, a total of 150 patients had their first PD catheter inserted at our University hospital. Table-1 depicts the demographic characteristics of the patients. There were 80 (53.3%) male and 70 (46.7%) female patients. The mean±SD age was 52.2±18.7 years. Mean±SD duration of APD was 15.4±5.8 months. Table-2 represents the technical complications during the study period. Assessment of PD catheter-related non-infectious complications showed that 10 out of 150 (6.7%) catheters were leaked in two weeks, late leakage occurred in 3 (2.0%) cases, hernias developed in 4 (2.7%), and transient catheter obstruction in 6 (4.0%) cases. Holding PD temporarily was sufficient to handle early leakage while late catheter leakage was treated surgically. Catheter obstruction was managed with t-PA (2 mg in 40 mL normal saline) instilled into the catheter. Thrombolysis was effective in all 6 cases, in 2 of them, occlusion occurred in the setting of acute peritonitis. One patient developed a leak at the catheter exit site within 24 hours after treatment. No recorded intraperitoneal bleeding and no changes were observed in systemic coagulation indices i.e. prothrombin time (PT), activated partial thromboplastin time (aPTT), fibrinogen degradation products (FDP), and fibrinogen levels. None of our patient had catheter migration.

There was no recording of bowel perforation or serious hemorrhage in the study population (Table 2). A total of 25
(17.3%) catheters had to be removed (Table 3). Indications for catheter removal were successful transplantation (n = 7), hernia (n = 4), catheter malfunction (n = 3; 2 ultrafiltration failure and 1 omental wrapping), unresolved peritonitis (n = 3), severe tunnel infection (n = 3), abdominal surgery (n = 1) and psychological causes (n = 4), while catheter replacement was performed in 3 (2.7%) because of omental capture (n = 1), and ultrafiltration failure (n = 2). As for infectious complications; peritonitis occurred in 20 (13.3%), ESI in 21 (14.0%), and tunnel infection in 4 (2.7%) patients. *Staphylococcus epidermidis* was the most common infectious agent and was isolated in 7 (4.7%) cases; followed by *Staphylococcus aureus* in 6 (4.0%) cases, *E. coli* in 3 (2.0%) cases and *Klebsiella species* in 2 (1.3%). Negative cultures were seen in 2 (1.3%) cases. Peritonitis responded well to medical treatment in the majority of cases (n = 17; 85%), while it was unresolved in 3 cases. During the study period, 10 (6.7%) patients were transferred to hemodialysis (4 with hernia, 3 with severe tunnel infection and 3 because of unresolved peritonitis), 7 (4.0%) patients received renal transplantation and 5 (3.3%) died (Figure 3). Causes of death were acute pulmonary embolism in 1, severe bilateral pneumonia in 1 and acute myocardial infarction in 3 patients. The study lasted 36 months with a mean APD duration of 15.4±5.8 months. Excluding patients who had successful renal transplantation (n = 7), abdominal surgery (n = 1) and catheter removal because of psychological problems (n = 4), catheter survival was 91.3% (Figure 4).

**Table 2:** Technical complications at the end of study.

| Complication       | n(%) |
|--------------------|------|
| Bowel perforation  | 0 (0.0) |
| Hemorrhage*        | 0 (0.0) |
| Ultrafiltration failure | 2 (1.3) |
| Omental wrapping   | 1 (0.7) |
| Catheter migration | 0 (0) |
| Early leakage**    | 10 (6.7) |
| Late leakage       | 3 (2.0) |
| Transient obstruction | 6 (4.0) |
| Catheter replacement | 3 (2.0) |

*Hemorrhage in the rectus muscle or pelvic cavity

**Within two weeks

**Table 3:** Causes of catheter removal during the study period.

| Cause                        | n(%) |
|------------------------------|------|
| Successful kidney transplant | 7 (4.7%) |
| Hernia                       | 4 (2.7%) |
| Ultrafiltration failure      | 2 (1.3%) |
| Omental wrapping             | 1 (0.7%) |
| Psychological                | 4 (2.7%) |
| Abdominal surgery            | 1 (0.7%) |
| Severe tunnel infection      | 3 (2.0%) |
| Unresolved peritonitis       | 3 (2.0%) |
| Total number                 | 25 (16.8%) |
| Technical survival*          | 137 (91.3%) |

*Technical survival after excluding catheter removal because of kidney transplant, abdominal surgery and psychological causes.

Discussion

This observational study aimed at evaluating the safety and efficacy of our new three-cuff PD catheter with low-entry
technique. In a previous publication, we reported an 18-month experience with our new catheter vs. the conventional Tenckhoff catheter [17]. The three-cuff PD catheter had zero rate of catheter migration or displacement. In our report we demonstrated that dialysis adequacy improved with the new catheter because of short fill and drainage times. In addition, the new catheter survival exceeded the ISPD recommendation. From December 2012, our new catheter was used for all incident PD patients [17].

Tenckhoff catheters have been the most widely used since the 1970s. Despite improvement in catheter survival rate over the last few years, complications such as catheter displacement erupted as one of the major causes of peritoneal dialysis technique failure [1,2]. Displacement or migration is often accompanied by obstruction of the holes in the catheter by omentum, and it usually impairs dialysis because of inadequate filling or drainage [18-22] and even cessation of outflow [22]. Catheter migration has been responsible for the frequently reported morbidity that necessitated catheter removal [23]. To restore the PD catheter function non-surgical options were tried and surgical maneuvers, with its negative impact on morbidity, were necessary in 85-90% of cases in some reports [22,24].

To overcome these problems, various catheter designs and insertion techniques have been described but the number of the concerned prospective and randomized studies is quite small [25]. When migration occurs, diathermy can be infused but drainage of the fluid from the peritoneal cavity is difficult. Suture fixation to the possible presence of ferrous impurities in the tungsten resonance (NMR) on patients with the self-locating catheter due to the presence of ferrous impurities in the tungsten catheter tip [33].

In addition, despite its promising results, the self-locating catheter is not used frequently in most of European countries. The reasons are unclear. Cost considerations might be important, with costs amounting to approximately US$310; the self-locating catheter is about 70% more expensive than the simple straight catheter described by Di Paolo, et al. in 1996 [32]. They reported no dislocations with the self-locating catheters, whereas nine dislocations occurred in control patients with the conventional Tenckhoff catheter (p =0.0003). In their report, however, there were no significant differences with respect to control for cuff extrusion, exit-site infections, leakage and peritoneal infection. In comparison, our catheter has zero cuff extrusion and a quite low incidence of peritoneal infection. Later, concerns were raised about the risk of carrying out abdominal nuclear magnetic resonance (NMR) on patients with the self-locating catheter due to the possible presence of ferrous impurities in the tungsten catheter tip [33].

Previous studies compared PD catheter survival in various catheter configurations. These studies included single-cuff and double-cuff, straight-end and curled-end catheters, and showed an incidence of catheter loss up to 35% depending on the catheter type [43]. Most of these studies, however, had a small patient sample or no control group. Outflow (one-way) is the most common type of PD catheter obstruction. This obstruction is caused by the closeness of the distal portion of the catheter to the omentum or intestine, which allows infusion of the solution, but little-to-no outflow. Persistent obstruction may require catheter manipulation with repositioning or replacement in many cases. Surgical laparotomy or laparoscopic epiploectomy of the greater omentum and epiploic appendices has been used to salvage a dysfunctional catheter [44]. Gadallah et al. [45] showed a lower catheter malfunction rate of 8.3% (including migration and omental capture), but those complications occurred within a shorter period of 2 weeks as compared with only 2.0% after a significantly longer duration in our series. With our insertion technique, the distal portion of the catheter being far away from the omentum and the intestine, which allows infusion of the solution, but little-to-no outflow. Persistent obstruction may require catheter manipulation with repositioning or replacement in many cases. Surgical laparotomy or laparoscopic epiploectomy of the greater omentum and epiploic appendices has been used to salvage a dysfunctional catheter [44]. Gadallah et al. [45] showed a lower catheter malfunction rate of 8.3% (including migration and omental capture), but those complications occurred within a shorter period of 2 weeks as compared with only 2.0% after a significantly longer duration in our series. With our insertion technique, the distal portion of the catheter being far away from the intestine and the omentum may explain the significantly lower incidence of omental wrapping and catheter malfunction. Our results confirmed that, as omental wrapping occurred only in 1 (0.7%) case.

Catheter related infection is the leading cause of peritoneal dialysis technique failure. In (CANUSA) study [46] peritonitis accounted for 15%-35% of hospital admissions. It was the major cause (40%-45%) of transfer to hemodialysis, and was associated with 7%-10% of deaths in PD patients. Recent data from Australia and Scotland [47,48] reported peritonitis rate of 0.6 episodes per year at risk or 1 episode every 20 months. According to ISPD
guidelines 2016 the overall peritonitis rate should be no more than 0.5 episodes per year equivalent to 1 per 24 patient-months [49] So far the best rate achieved was as low as 0.18 to 0.20 per year [50-52]. Peritonitis rate in our center prior to our current trial catheter was 0.2% equivalent to 1 episode per 60 patient-months. With our new 3-cuff PD catheter and new technique the rate of peritonitis was as low as 0.106 per patient-year equivalent to 1 episode of peritonitis per 112 patient-months i.e. far exceeding the ISPD recommendations and superior to the previous reports. This may probably be attributed to the presence of three cuffs which act as three barriers against peritoneal infections. The short intra abdominal catheter segment favorably influenced the infection outcome. One drawback of our new catheter that it is removal requires two incisions instead of one (compared with the conventional Tenckhoff catheter).

The present study showed that patients with the new catheter had fewer episodes of peritonitis, tunnel infection, cuff extrusion, catheter malfunction, obstruction, and leakage and without catheter displacement. Together, these factors had its positive impact on catheter survival as proved by our long-term study. In our opinion, our results are logical consequence of maintaining the correct position of the catheter tip in the Douglas cavity. If the tip remains in the bottom of this cavity, catheter displacement and cuff extrusion are unlikely, tunnel infection decreases, good catheter function is conserved (omentum is absent from the Douglas cavity), and therefore episodes of peritonitis subsequently decrease. We also believe that a larger patient population needs to be studied in a multicenter setting to confirm our findings.

Conclusion

Compared with other PD catheters, the new triple-cuff PD catheter demonstrated zero rate of catheter migration. Catheter survival and peritonitis rate far exceeded the ISPD recommendations, and other PD centers. We believe that our triple-cuff PD catheter with deep located short intra abdominal segment can help to minimize catheter complications, particularly migration and peritonitis. Our data need to be confirmed in larger prospective randomized trials comparing it with other modern catheter designs.

Acknowledgment

The authors thank all the PD Unit staff at King Fahd Hospital of the University. The authors also extend their appreciation to the staff and technicians of the laparoscopy unit for their remarkable help and support during the preparation of this study.

Conflict of Interest

The authors have no relationship with pharmaceutical companies or other entities such as employment contracts, consultancy, advisory boards, speaker bureaus, membership of Board Directors, stock ownership that could be perceived to represent a financial conflict of interest.

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