

ERKNet/ESPN

Workshop on fundamentals in pediatric dialysis

21 - 22 October 2021



Peritonitis and Other Complications of Peritoneal Dialysis in Children

Sevcan A. Bakkaloğlu

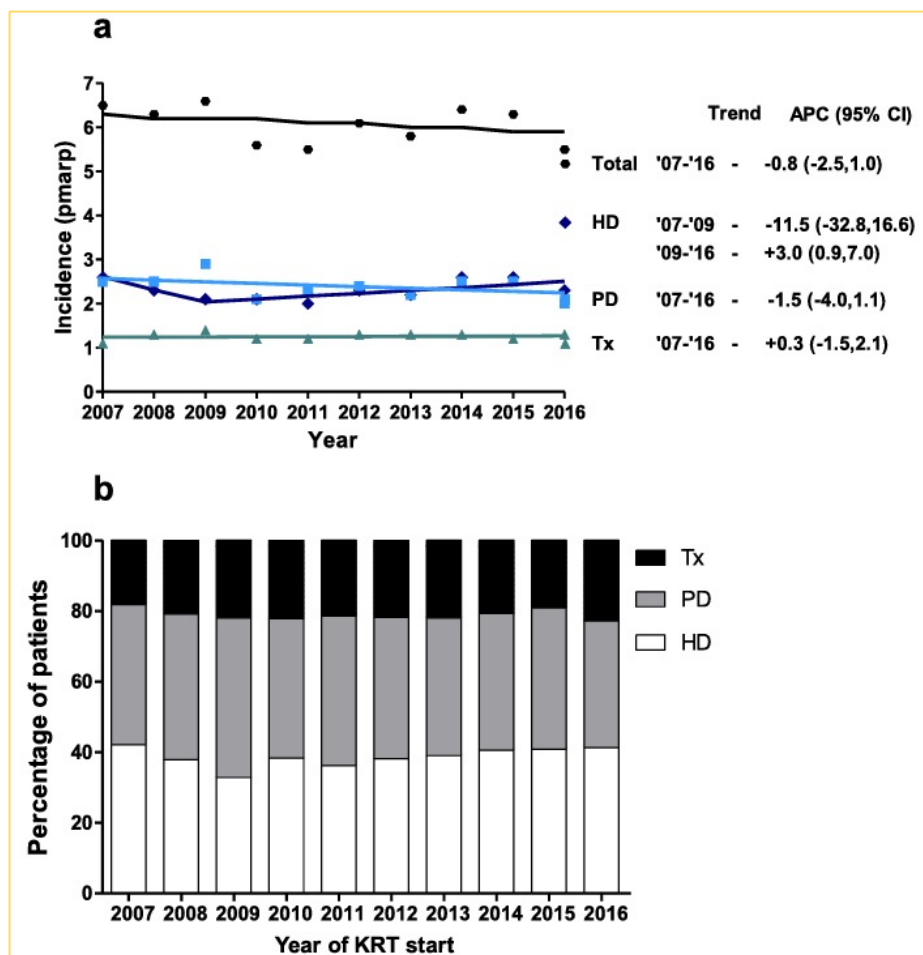
Gazi University School of Medicine

Department of Pediatric Nephrology

Ankara, TURKEY

Ten-year trends in epidemiology and outcomes of pediatric kidney replacement therapy in Europe: data from the ESPN/ERA-EDTA Registry

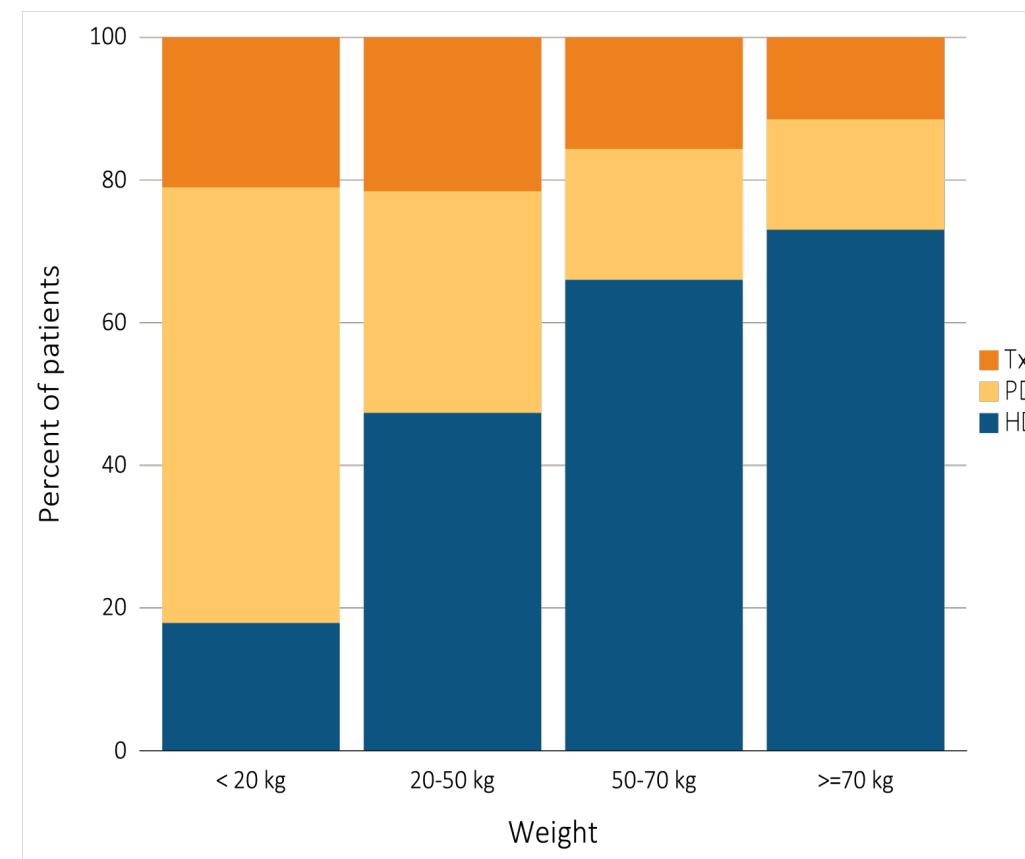
Marjolein Bonthuis¹ • Enrico Vidal² • Anna Bjerre³ • Özlem Aydoğ⁴ • Sergey Baiko⁵ • Liliana Garneata⁶ • Isabella Guzzo⁷ • James G. Heaf⁸ • Timo Jahnukainen⁹ • Marc Lilien¹⁰ • Tamara Mallett^{11,12} • Gabriel Mirescu^{13,14} • Elena A. Mochanova¹⁵ • Eva Nüsken¹⁶ • Katherine Rascher¹⁷ • Dimitar Roussinov¹⁸ • Maria Szczepanska¹⁹ • Michel Tsimaratos²⁰ • Askiti Varvara²¹ • Enrico Verrina²² • Bojana Veselinović²³ • Kitty J. Jager¹ • Jérôme Harambat²⁴



Pediatr Nephrol 2021

Trends in pediatric ESRD modality at initiation, by patient weight, 1996-2014

USRDS 2016 report



Complications of PD

- **Infectious complications**

- Peritonitis
- Catheter-related infections
 - Exit-site infections
 - Tunnel infections

- **Noninfectious complications**

- Mechanical
 - Leakage
 - Catheter migration
 - Cuff extrusion
 - Outflow obstruction
 - Hernias

Adequacy and ultrafiltration problems

- Inadequate solute clearance
- Poor compliance
- Decreased peritoneal permeability
- Inadequate ultrafiltration
- Fast transport status
- Encapsulated peritoneal sclerosis

Metabolic complications

- Hyperglycemia
- Hyperinsulinemia
- Hypertriglyceridemia
- Hypokalemia
- Magnesium alterations

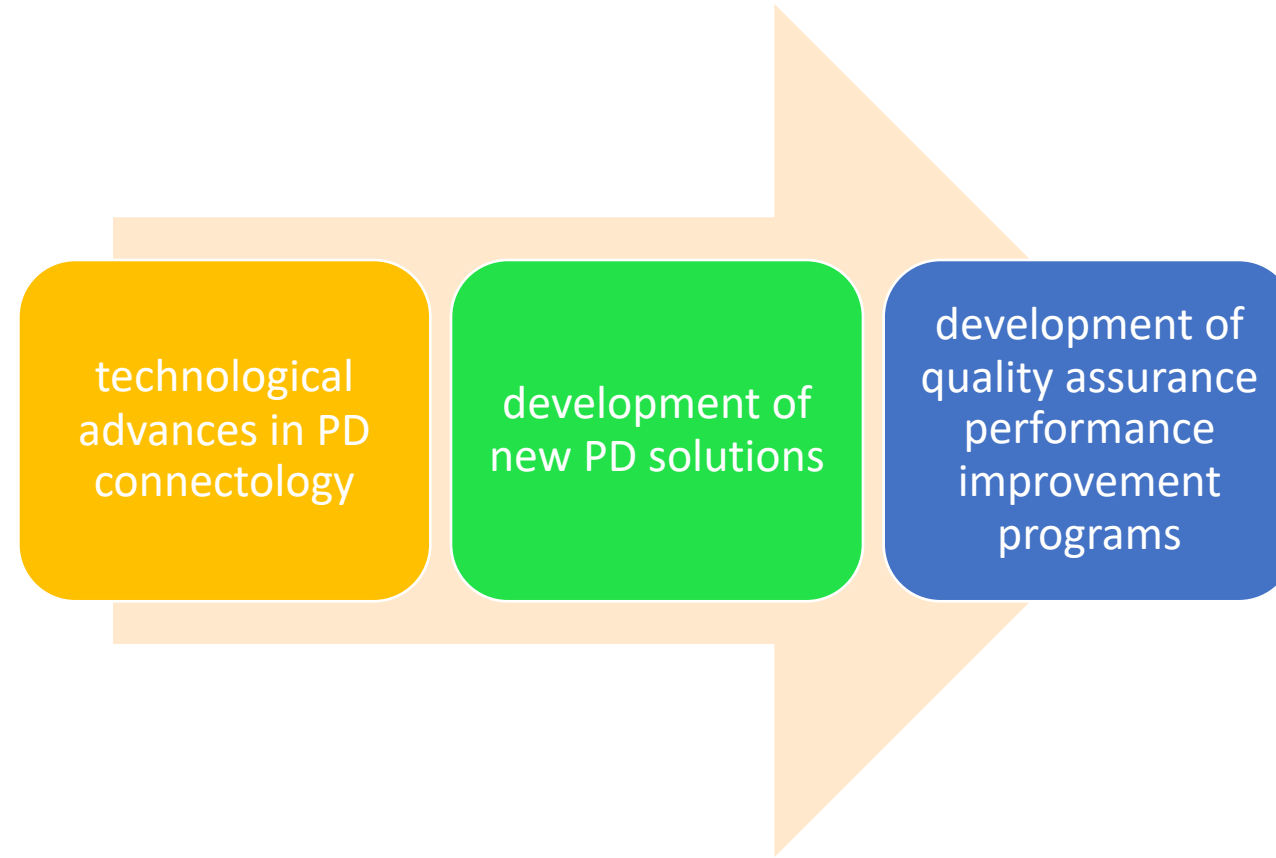
- **Other complications**

Hemoperitoneum

Pneumoperitoneum

Pancreatitis

Ischemic colitis and necrotizing enterocolitis



Peritonitis remains the
most common
complication of CPD

Scan me!



IPPN



NAPRTCS



USRDS

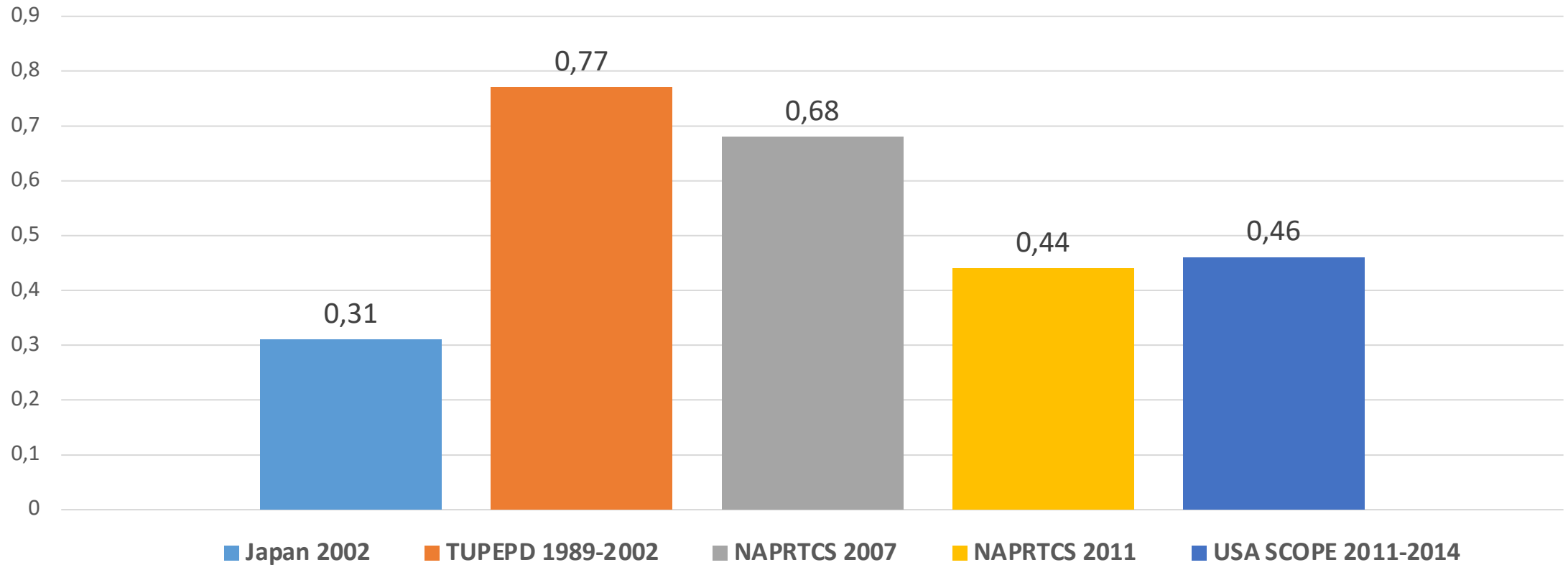


SCOPE



TSPN RRT

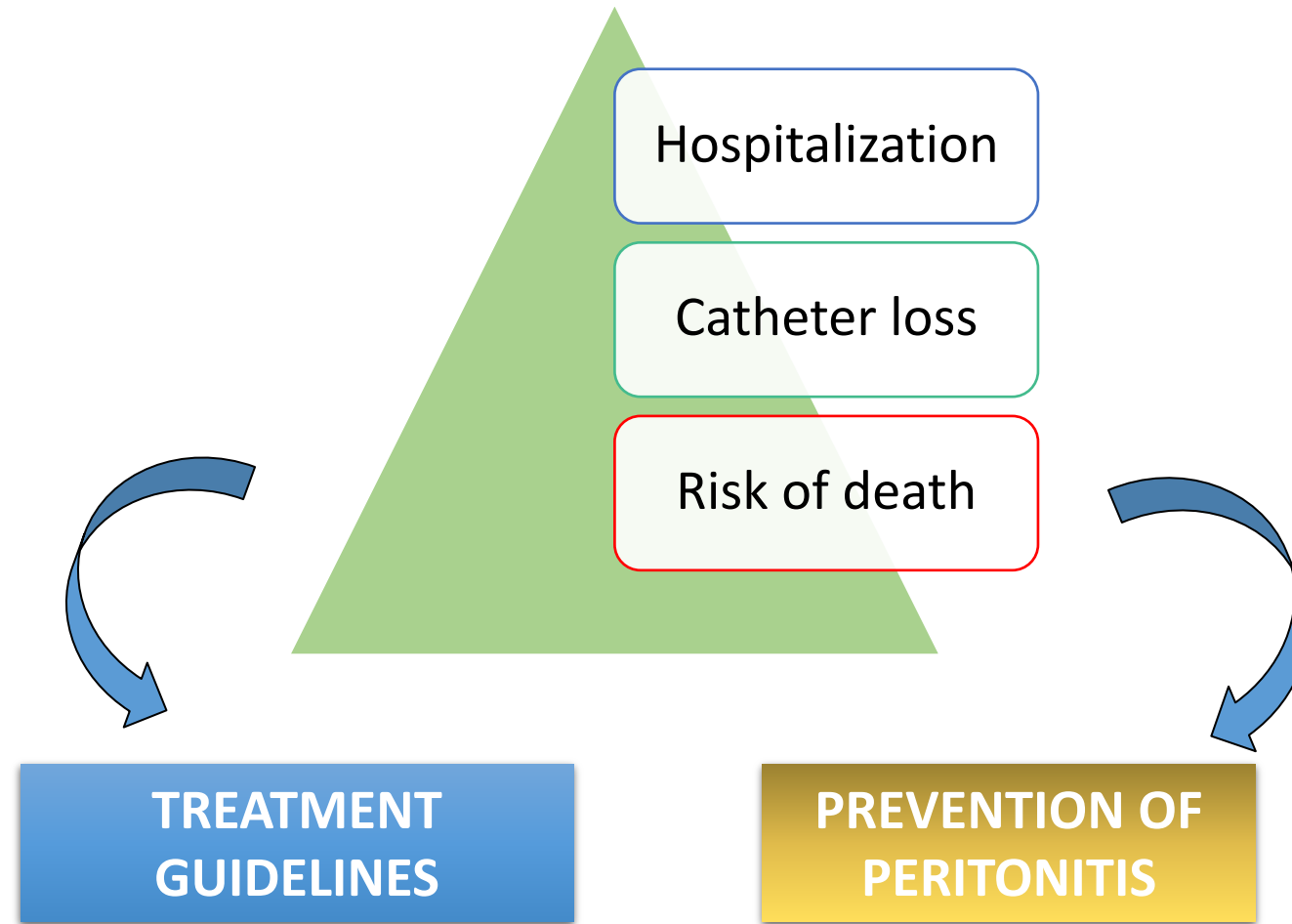
Peritonitis rate episode per year



NAPRTCS 2011 - Significant improvement is seen since 2002 with the annualized rate of infection **decreasing from 0.79 in 1992-1996 to 0.44 in recent years.**

NAPRTCS 2007, Honda M, Proc Pediatr PD Conf 2002, Akman S, Pediatr Int 2009 ,
NAPRTCS 2011, Setha J, CJASN 2017

Peritonitis



Prevention of Peritonitis

Catheter related interventions to reduce peritonitis risk

- Use double-cuff catheters

- Lateral/downward subcutaneous tunnel configuration

Antimicrobial prophylaxis

- Preoperative antibiotics at catheter insertion

- Exit site antimicrobial application

- Fungal prophylaxis

Prevention of contamination

- Experienced nursing personnel

- Long training period, retraining, home visits

- Avoidance of spiking technology

- Contamination protocols

Development of continuous quality improvement programs

- Tracking and questioning infectious episodes as well as causative agent

- Monthly team meetings (evaluation of root cause; plan to prevention of recurr.)

- Reevaluation of protocols of the PD program

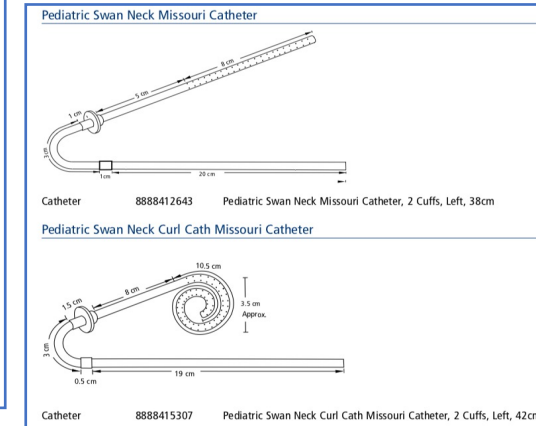
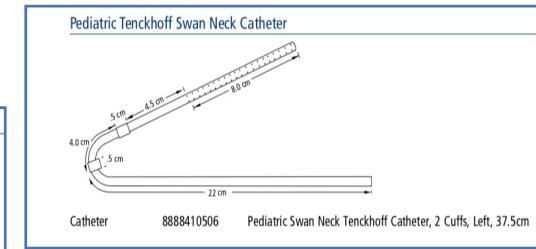
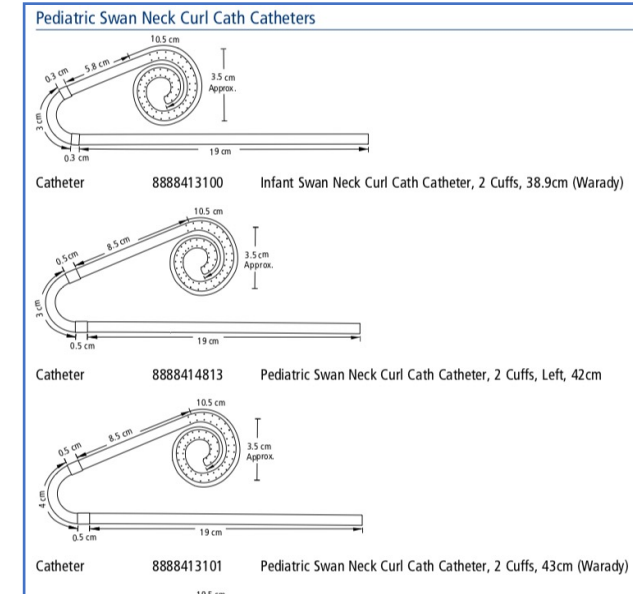
Prevention of Peritonitis

Catheter related interventions to reduce peritonitis risk

Use double-cuff catheters

Lateral/downward subcutaneous tunnel configuration

- Catheter selection
 - Straight vs coiled
 - Single cuff vs double cuff
 - Preformed curve (swan-neck) or straight tunnel
- Exit-site orientation
 - Lateral or downward
 - Subcutaneous burying
- Catheter insertion techniques
 - Blind percutaneous
 - Open surgical
 - Laparoscopic
 - Minilaparoscopic
 - Peritoneoscopic single trocar



No firm agreement on exact catheter type and configuration

No technique of PD catheter implantation that has consistently proven to be superior for the prevention of peritonitis

PD CATHETER SIZE

CENTER's SURGICAL EXPERTISE in the APPLIED METHOD

Locate superficial cuff 2 cm from the exit site

Use the smallest exit hole

No incision, no sutures at the exit site

Catheter anchoring and immobilization

Dressing changes should be avoided in the first week, then weekly up to 6 wk

If possible, do not use the catheter at least for two weeks

Prevention of Peritonitis

Catheter related interventions to reduce peritonitis risk

Use double-cuff catheters

Lateral/downward subcutaneous tunnel configuration

Antimicrobial prophylaxis

Preoperative antibiotics at catheter insertion

Exit site antimicrobial application

Fungal prophylaxis

Risk factors for *Pseudomonas* peritonitis

Use of saline or soap for cleansing

Exit site care > twice per week

Use of exit site mupirocin

Schaefer et al. Kidney Int 2007

ES Cleansing solutions

Povidone-iodin

Chlorhexidine solution

Amuchina solution/hypochlorite solution)

Topical antimicrobials

Mupirocin cream

Gentamicin cream or ointment

Ciprofloxacin otologic solution

Antibacterial honey

Polysporin triple ointment

ISPD 2016 Adult guideline

- Daily topical application of antibiotic (mupirocin or gentamicin) cream or ointment to the catheter exit site is recommended
- Mupirocin resistance has been reported, particularly with intermittent use but not daily use



Bernardini J

Prevention of Peritonitis

Catheter related interventions to reduce peritonitis risk

Use double-cuff catheters
Lateral/downward subcutaneous tunnel configuration

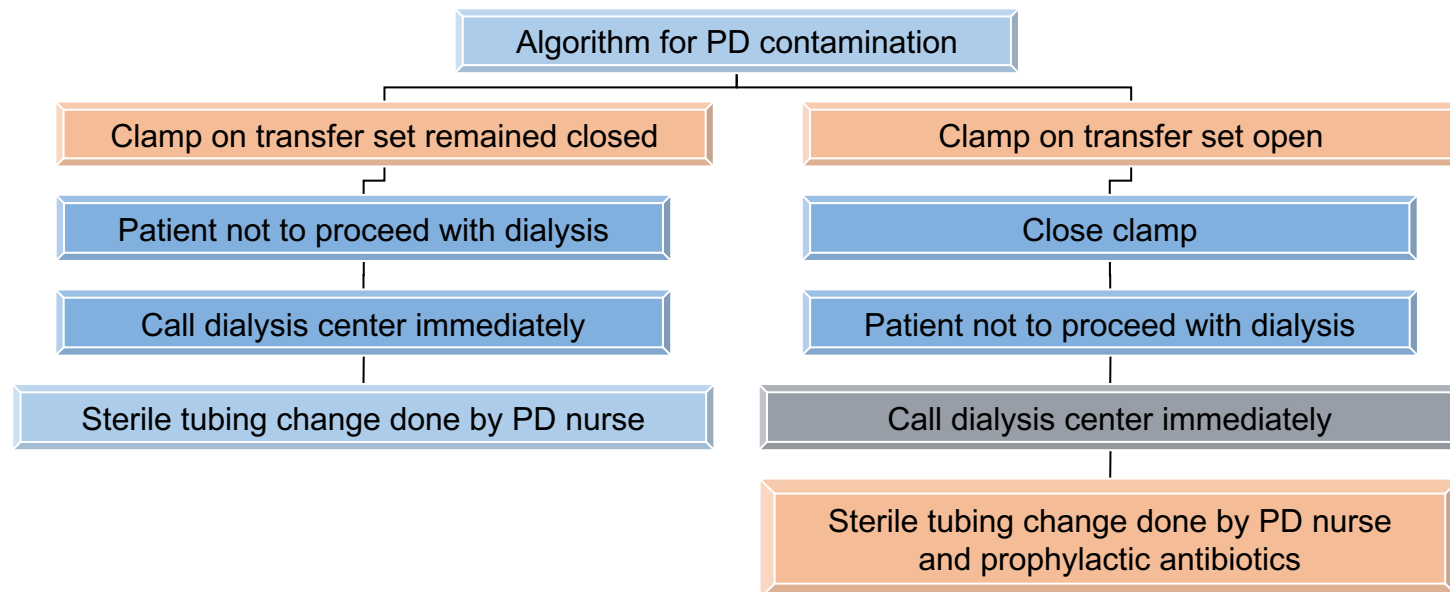
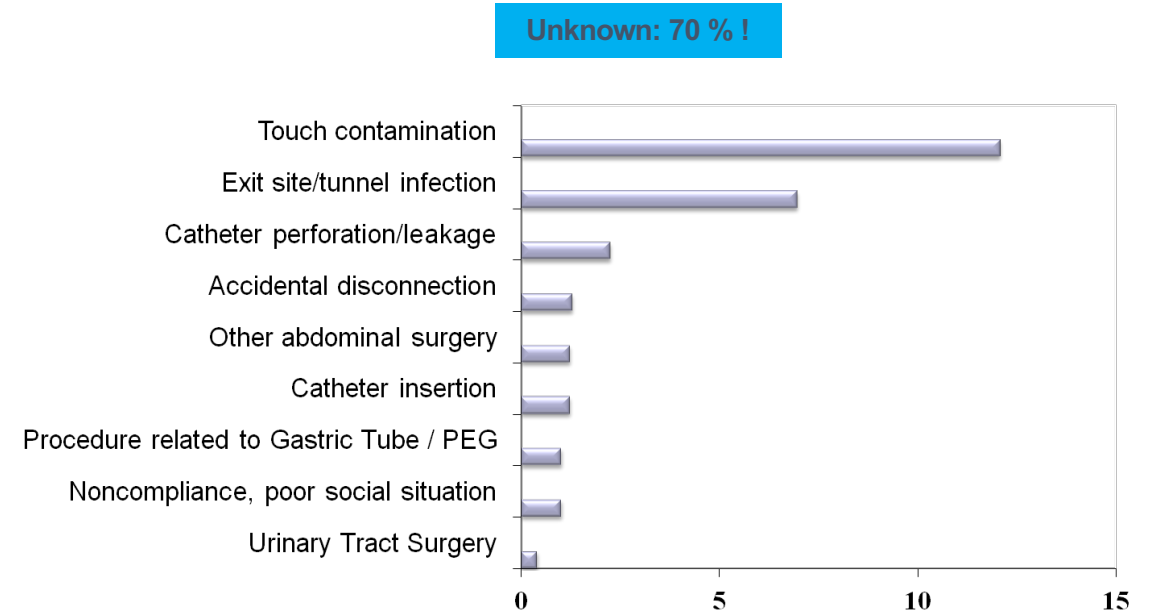
Antibiotic prophylaxis

Preoperative antibiotics at catheter insertion
Exit site antimicrobial application
Fungal prophylaxis

Prevention of contamination

Experienced nursing personnel
Long training period, retraining, home visits
Avoidance of spiking technology
Contamination protocols

Peritonitis: Source of Infection



Prevention of Peritonitis

Catheter related interventions to reduce peritonitis risk

Use double-cuff catheters

Lateral/downward subcutaneous tunnel configuration

Antibiotic prophylaxis

Preoperative antibiotics at catheter insertion

Exit site antimicrobial application

Fungal prophylaxis

Prevention of contamination

Experienced nursing personnel

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Development of continuous quality improvement programs

Tracking and questioning infectious episodes as well as causative agent

Monthly team meetings (evaluation of root cause; plan to prevention of recurr.)

Reevaluation of protocols of the PD program

SCOPE Collaborative – Standardized Care to Improve Outcomes in Pediatric End Stage Kidney Diseases

- › Collaboratives
- › Pediatric Quality Policy and Measures
- › Partnerships
- › Pediatric Quality Award
- › Legacy Quality Programs
- › Data Analytics & Research
- › Peer Networking & Collaboration
- › Pediatric Learning Solutions
- › Group Purchasing & Supply Chain
- › Insurance Services



The Standardized Care to Improve Outcomes in Pediatric Endstage Renal Disease (SCOPE) collaborative prevents infections in pediatric peritoneal dialysis and hemodialysis patients using large-scale collaboration to identify and spread effective interventions across pediatric care settings.

Focus

Peritoneal Dialysis (PD) – Preventing infections in PD patients reduces hospitalizations and allows patients to continue home-based dialysis and maintain normal school and family activities.

Hemodialysis (HD) – Similarly, preventing infections in HD patients reduces hospitalizations and the need for additional invasive procedures. Additionally, SCOPE is contributing to new knowledge about children on dialysis while improving outcomes.

✉ Cherie Thomas
913-981-4199

✉ Megan Esporas
919-484-9890

Scope Resources

- › SCOPE Flyer
- › SCOPE Teams
- › SCOPE FAQs
- › SCOPE Publications and Presentations

Scope Additional Resources

- › SCOPE Enrollment Brief
- › Enrollment Form
- › SCOPE Charter
- › SCOPE HD Charter
- › SCOPE Enrollment Toolkit

Impact

(as of July 2021)

PD - peritoneal dialysis; HD - hemodialysis



430 PD-related hospitalizations prevented



733 peritoneal infections prevented



Over \$12.9 million saved hospitalization costs for PD-related infections and

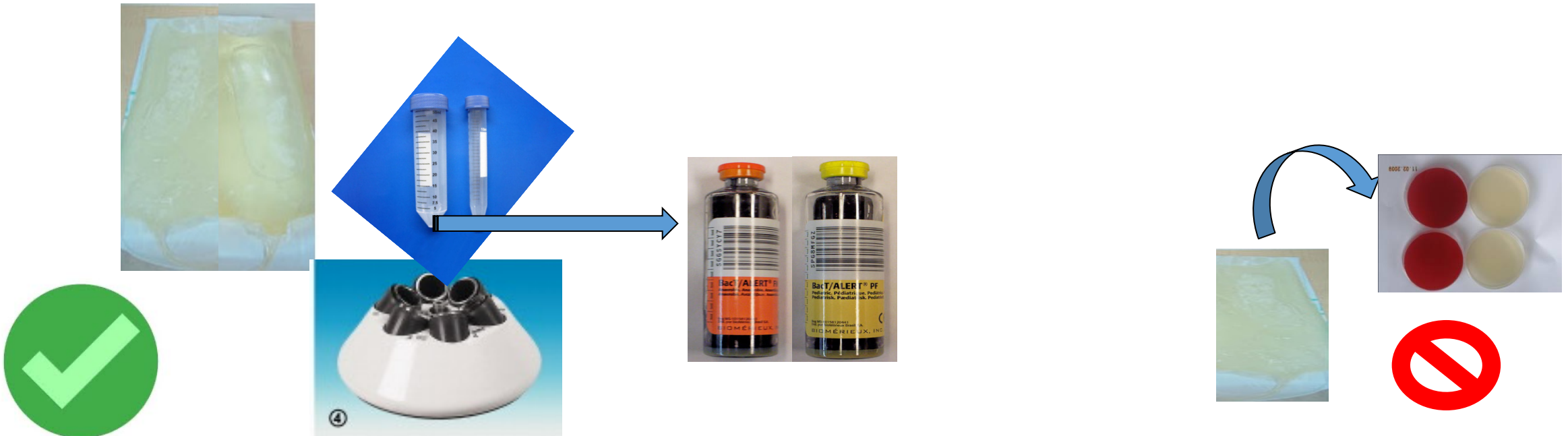


CDC Partnership



Peritonitis

- cell count
 - differential count
 - culture to confirm the diagnosis of peritonitis
- WBC > 100/mm³, and at least 50% of the WBCs are PMNL
 - centrifugation of PD effluent
 - culture of sediment
 - blood-culture bottles as the standard culture technique



Spectrum of Causative Organisms

| | IPPR (501 episode) | | IPPN (1456 episode) | | SCOPE (389 episode) |
|---------------|-----------------------|----|------------------------|-----|------------------------|
| Gram (+) | 44% | | 63% | | 38% |
| Gram (-) | 25% | | 33% | | 20% |
| Sterile | 31% | | | 33% | 25% |
| Fungal | | 2% | 4% | | 8% |
| Polymicrobial | | | | | 10% |

| | |
|-----------------|-----|
| CNS | 25% |
| S. aureus | 22% |
| E.coli | 8% |
| Streptococci | 7% |
| Pseudomonas sp. | 6% |
| Enterococci | 6% |

Geographic variability

Environmental – Humidity, climate

PD practice – ES care, topical antibiotic prophylaxis

Peritoneal Dialysis International

Peritoneal Dialysis International, Vol. 32, pp. 532-586
doi: 10.3747/pdi.2011.00091

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ISPD GUIDELINES/RECOMMENDATIONS

CONSENSUS GUIDELINES FOR THE PREVENTION AND TREATMENT OF
CATHETER-RELATED INFECTIONS AND PERITONITIS IN PEDIATRIC
PATIENTS RECEIVING PERITONEAL DIALYSIS: 2012 UPDATE

Bradley A. Warady,¹ Sevcn Bakkaloglu,² Jason Newland,¹ Michelle Cantwell,³ Enrico Verrina,⁴ Alicia Neu,⁵
Vimal Chadha,¹ Hui-Kim Yap,⁶ and Franz Schaefer⁷

Peritoneal Dialysis International

Peritoneal Dialysis International, Vol. 36, pp. 481-508
www.PDIConnect.com

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ISPD GUIDELINES/RECOMMENDATIONS

ISPD PERITONITIS RECOMMENDATIONS: 2016 UPDATE ON PREVENTION AND TREATMENT

Philip Kam-Tao Li,¹ Cheuk Chun Szeto,¹ Beth Piraino,² Javier de Arteaga,³ Stanley Fan,⁴ Ana E. Figueiredo,⁵
Douglas N. Fish,⁶ Eric Goffin,⁷ Yong-Lim Kim,⁸ William Salzer,⁹ Dirk G. Struijk,¹⁰
Isaac Teitelbaum,¹¹ and David W. Johnson¹²

Peritoneal Dialysis International

Peritoneal Dialysis International, Vol. 37, pp. 141-154
www.PDIConnect.com

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ISPD GUIDELINES/RECOMMENDATIONS

ISPD CATHETER-RELATED INFECTION RECOMMENDATIONS: 2017 UPDATE

Cheuk-Chun Szeto,¹ Philip Kam-Tao Li,¹ David W. Johnson,² Judith Bernardini,³ Jie Dong,⁴ Ana E. Figueiredo,⁵
Yasuhiko Ito,⁶ Rumez Kazancioglu,⁷ Thyago Moraes,⁸ Sadie Van Esch,⁹ and Edwina A. Brown¹⁰

Peritoneal Dialysis International

Peritoneal Dialysis International, Vol. 31, pp. 614-630
doi: 10.3747/pdi.2011.00057

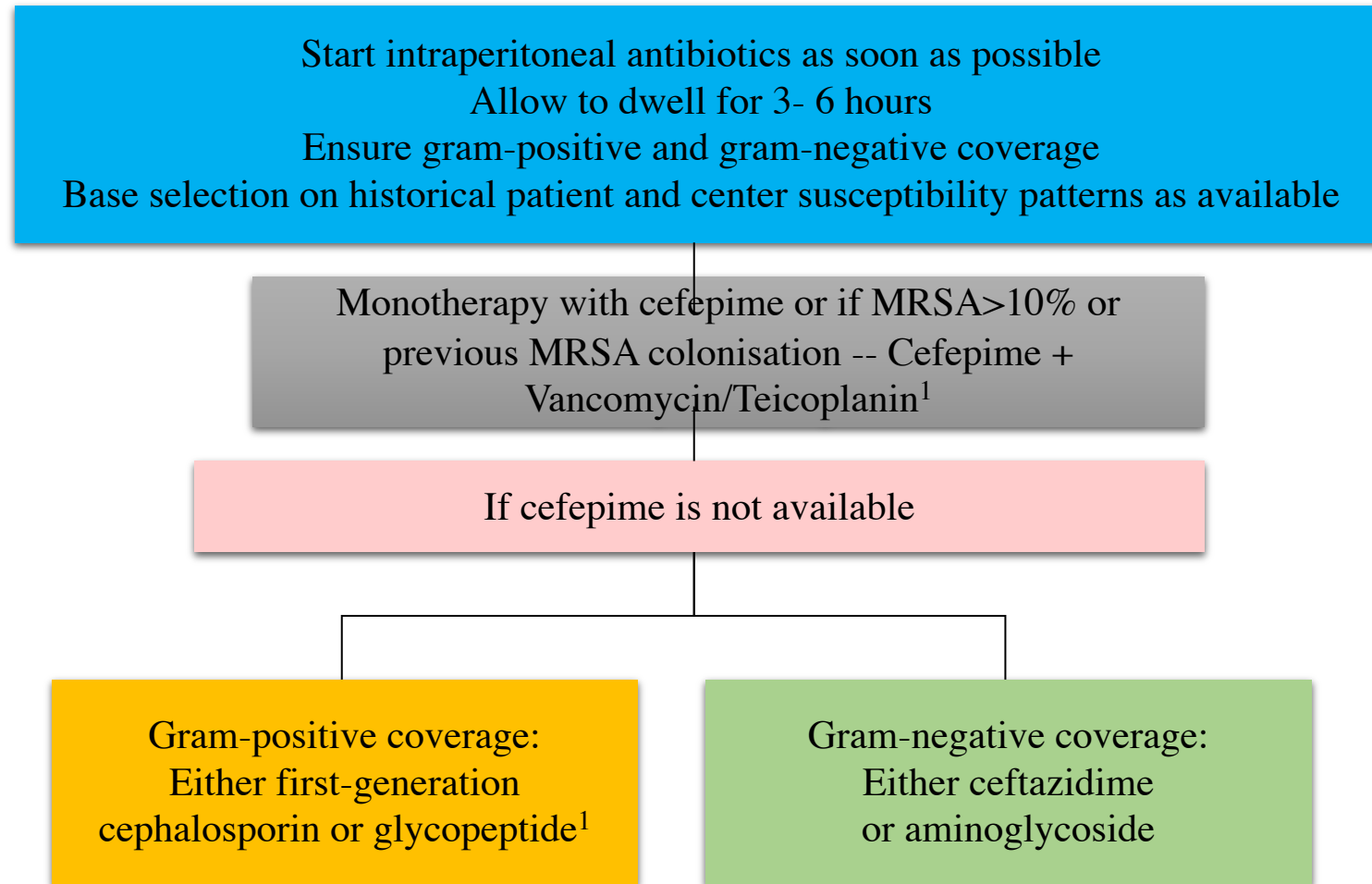
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SPECIAL ARTICLE

ISPD POSITION STATEMENT ON REDUCING THE RISKS OF PERITONEAL
DIALYSIS-RELATED INFECTIONS

Beth Piraino,¹ Judith Bernardini,¹ Edwina Brown,² Ana Figueiredo,³ David W. Johnson,⁴ Wai-Choong Lye,⁵
Valerie Price,⁶ Santhanam Ramalakshmi,⁷ and Cheuk-Chun Szeto⁸

Empiric antibiotic therapy



If the center's MRSA rate exceeds 10% or patient has history of MRSA colonization, glycopeptide (vancomycin / teicoplanin) should be added to cefepime or should replace the first generation cephalosporin for gram-positive coverage. Glycopeptide usage can also be considered if patient has a history of severe allergy to penicillins and cephalosporins.

| | Continuous therapy | | Intermittent therapy |
|------------------------------|---|------------------|---|
| | Loading dose | Maintenance dose | |
| Aminoglycosides ^a | | | |
| Gentamicin | 8 mg/L | 4 mg/L | anuric: 0.6 mg/kg non-anuric: 0.75 mg/kg. |
| Netilmycin | 8 mg/L | 4 mg/L | |
| Tobramycin | 8 mg/L | 4 mg/L | |
| Cephalosporins | | | |
| Cefazolin | 500 mg/L | 125 mg/L | 20 mg/kg |
| Cefepime | 500 mg/L | 125 mg/L | 15 mg/kg |
| Cefotaxime | 500 mg/L | 250 mg/L | 30 mg/kg |
| Ceftazidime | 500 mg/L | 125 mg/L | 20 mg/kg |
| Glycopeptides ^b | | | |
| Vancomycin | 1000 mg/L | 25 mg/L | 30 mg/kg; repeat dosing 15 mg/kg every 3-5 days |
| Teicoplanin ^c | 400 mg/L | 20 mg/L | 15 mg/kg q 5 – 7 days |
| Penicillins ^a | | | |
| Ampicillin | ----- | 125 mg/L | ----- |
| Quinolones | | | |
| Ciprofloxacin | 50 mg/L | 25 mg/L | ----- |
| Others | | | |
| Aztreonam | 1000 mg/L | 250 mg/L | ----- |
| Clindamycin | 300 mg/L | 150 mg/L | ----- |
| Imipenem/Cilastin | 250 mg/L | 50 mg/L | ----- |
| Oral | | | |
| Linezolid | < 5 yrs: 30 mg/kg/day divided TID; 5-11 yrs: 20 mg/kg/day divided BID; ≥ 12 yrs 600 mg/dose BID | | |
| Metronidazole | 30 mg/kg/day divided TID | | |
| Rifampin | 10-20 mg/kg/day divided BID (max daily dose 600 mg) | | |
| Antifungals | | | |
| Fluconazole | 6 – 12 mg/kg IP, IV or PO every 24-48 hrs (max dose 400 mg) [#] | | |
| Caspofungin | IV only: initial dose 70 mg/m ² on day 1 (max dose 70 mg); Subsequent dosing 50 mg/m ² daily (max dose 50 mg) | | |

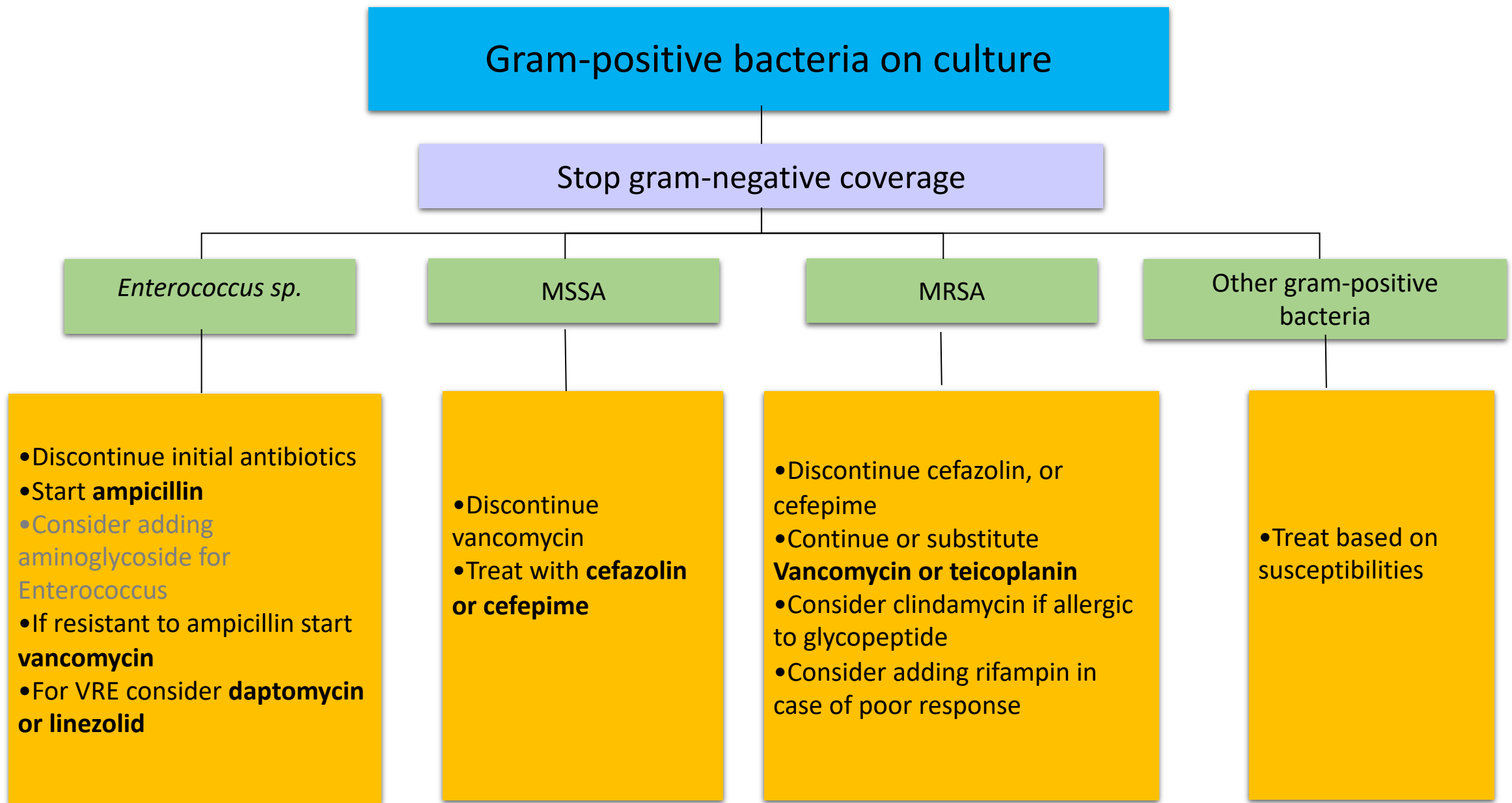


Figure 2. Gram-positive organism on culture. MRSA-methicillin resistant *S. aureus*; methicillin sensitive *S. aureus*; VRE-vancomycin resistant enterococci.

Gram-positive bacteria and the recommended antibiotics and length of therapy

| | Recommended Antibiotic(s)* | Length of Therapy |
|--|---|-------------------|
| Methicillin-resistant <i>S. aureus</i> | Vancomycin/Teicoplanin, Clindamycin | 3 weeks |
| Methicillin-susceptible <i>S. aureus</i> | Cefazolin, Cefepime | 3 weeks |
| Coagulase negative staphylococci | Vancomycin/Teicoplanin, Clindamycin if MR | 2 weeks |
| Enterococcus sp. | Ampicillin, Vancomycin/Teicoplanin | 2-3 weeks |
| Vancomycin resistant Enterococcus | Daptomycin, Linezolid | 2-3 weeks |
| Streptococcus species | Ampicillin, Cefazoline, Cefepime | 2 weeks |

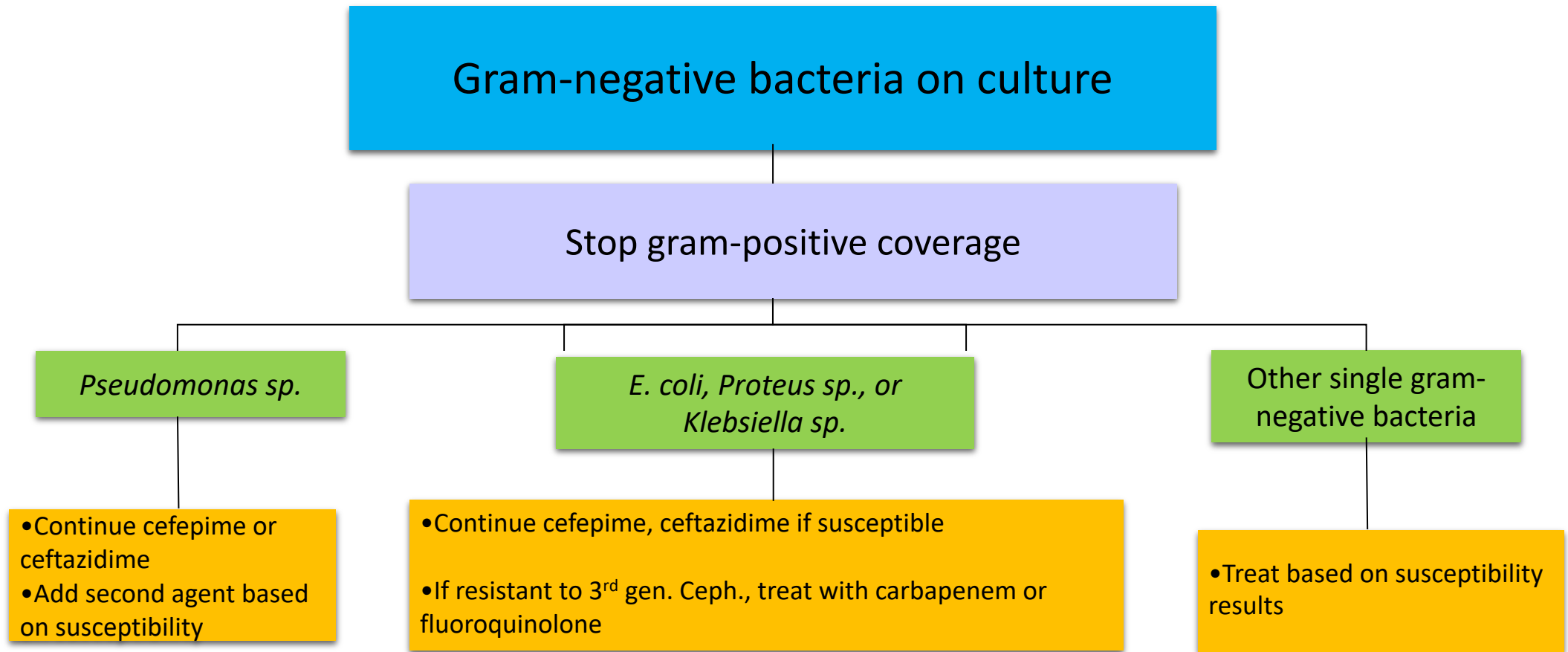


Figure 3. Gram-negative organism on culture


Gram-negative bacteria and the recommended antibiotics and length of therapy

| Bacteria | Recommended Antibiotic(s)* | Length of Therapy |
|---|--|-------------------|
| <i>E. coli</i> , <i>Klebsiella</i> sp. | Cefazolin, Cefepime, Ceftazidime, Ceftriaxone/ Cefotaxime | 2 weeks |
| <i>E. coli</i> , <i>Klebsiella</i> sp. resistant to 3 rd generation Cephalosporins | Carbapenem** or Fluoroquinolone | 3 weeks |
| <i>Enterobacter</i> sp., <i>Citrobacter</i> sp., <i>Serratia</i> sp., <i>Proteus</i> sp. | Cefepime, Ceftazidime or Carbapenem** | 2-3 weeks |
| <i>Acinetobacter</i> sp. | Cefepime, Ceftazidime or Carbapenem | 2-3 weeks |
| <i>Pseudomonas</i> species | Cefepime, Ceftazidime, Piperacillin or Ticarcillin, plus Aminoglycoside or Fluoroquinolone | 3 weeks |
| <i>Stenotrophomonas maltophilia</i> | Trimethoprim/Sulfamethoxazole, Ticarcillin/Clavulanic acid | 3 weeks |

Fungal peritonitis

- 2%-8% of all peritonitis episodes in children
- Risk factors
 - Prior antibiotic use
 - Gastrostomy ?
 - Antifungal prophylaxis during antibiotic usage in programs with high rates of fungal peritonitis
- If fungi are identified by Gram stain or culture of peritoneal effluent, therapy should consist of treatment with **an antifungal agent and early catheter removal**
- Following catheter removal, antimycotic therapy be administered for 2 weeks or longer following complete resolution of the clinical symptoms of infection

Treatment

- Fluconazole for Candida species
- Caspofungin for non responding non-albicans Candida
- Voriconazole for Aspergillus
- Treatment duration following catheter removal should be 2 weeks or longer following complete resolution of the clinical symptoms of infection
- Amphotericin B 
 - Poor peritoneal penetration
 - Intraperitoneal irritation and abdominal pain

Difficult peritonitis cases in children undergoing chronic peritoneal dialysis: relapsing, repeat, recurrent and zoonotic episodes

Sevcan A. Bakkaloglu · Bradley A. Warady



Relapsing peritonitis

Infectious locus poorly accessible to antibiotics (catheter tunnel, fibrin, biofilm)

IPPR - 2.5-fold risk of permanent PD technique failure

While waiting in vitro susceptibility results, reinitiation of empiric therapy with consideration of the susceptibilities of the original bacteria

Postempirically, choosing an antibiotic other than cefazolin
IP urokinase or t-PA

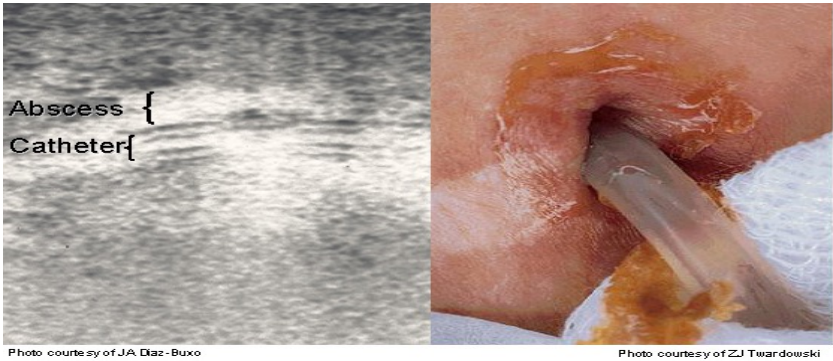
Remove the PD catheter as soon as peritonitis is controlled by antibiotic therapy in the setting of
relapsing peritonitis associated with persistent or recurrent tunnel infection
a second peritonitis relapse

Indications for catheter removal and replacement

| Catheter removal | | Reinsertion |
|--|--|--|
| | Refractory bacterial | After 2-3 days |
| | Fungal | After >2 weeks |
| | ESI/TI in peritonitis with the same organism (mainly, <i>S. aureus</i> and <i>P. aeruginosa</i> ; except CNS) | After 2-3 weeks |
| Simultaneous removal and replacement of the catheter | Relapsing or refractory ESI/TI (including <i>P. aeruginosa</i>) | |
| | Relapsing peritonitis | |
| Relative indications for removal | Repeat peritonitis | After 2-3 weeks |
| | Peritonitis with multiple enteric organisms due to an intra-abdominal pathology/ abscess; so-called surgical peritonitis | Dependent upon the clinical course of the patient ; at least 2-3 weeks |

Save the peritoneum
not save the catheter!

Exit-site / Tunnel infections

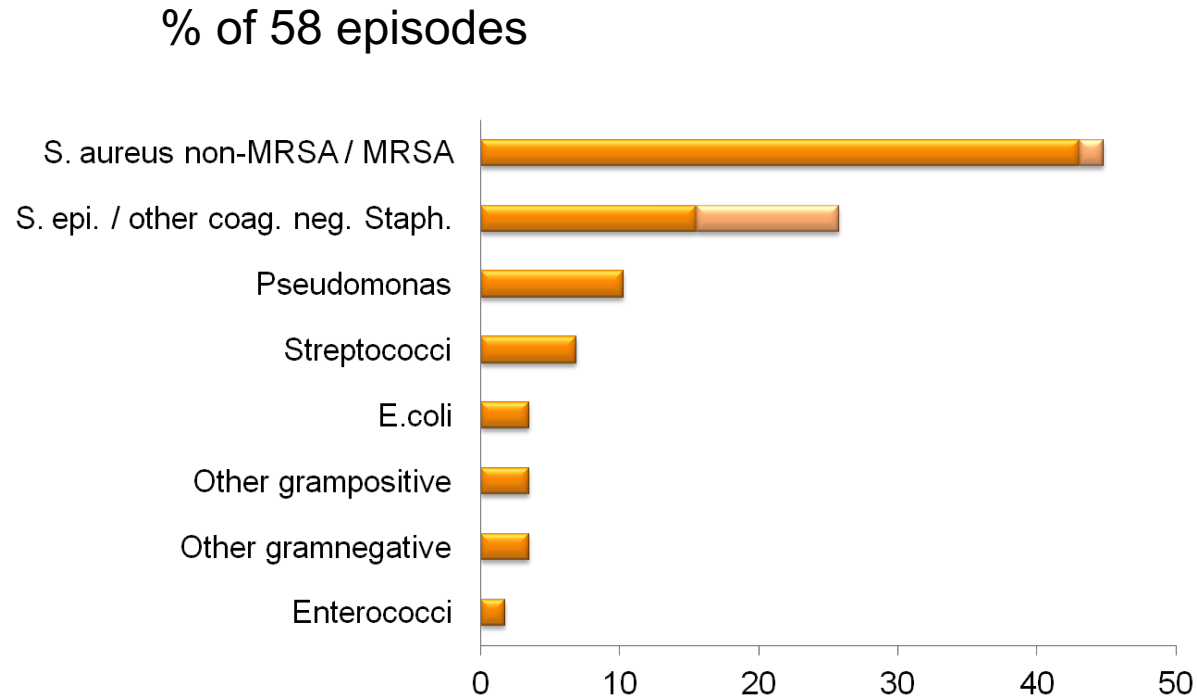


Exit-site scoring system

| | 0 Points | 1 Point | 2 Points |
|------------------|----------|---------------------|------------------------------------|
| Swelling | No | Exit only (< 0.5cm) | Including part of or entire tunnel |
| Crust | No | < 0.5cm | > 0.5cm |
| Redness | No | < 0.5cm | > 0.5cm |
| Pain on pressure | No | Slight | Severe |
| Secretion | No | Serous | Purulent |

For ESI exit-site score should be 4 or greater with or without positive culture (2 or greater if culture is positive)
 For TI exit-site score should be 6 or greater with or without purulent drainage

Causative Organisms at Exit Site



Catheter-related peritonitis - IPPR

2-fold risk of peritonitis treatment failure

3-fold risk of catheter exchange

Treatment of Exit-site / Tunnel Infections

- Exit-site infections:

- Oral antibiotic therapy when culture results and susceptibilities available
- Gram positive usually penicillinase-resistant penicillin or cefalexin
- Gram negative IP ceftazidime, combination therapy for *Pseudomonas*
- a minimum of 2 weeks (3 weeks for *S. aureus* and *P. aeruginosa*, max 4 weeks)
- at least 7 days following complete resolution of the infection

Tunnel infections:

- Antibiotic therapy after culture and susceptibility results have been obtained
- Signs of severe infection, and/or a history of *S. aureus* or *P. aeruginosa* initiation of empiric therapy should be considered
- Oral, intraperitoneal or intravenous routes
- MRSA IV Glycopeptide
- Treatment duration should be 2-4 weeks

Oral antibiotics used in exit-site and tunnel infection

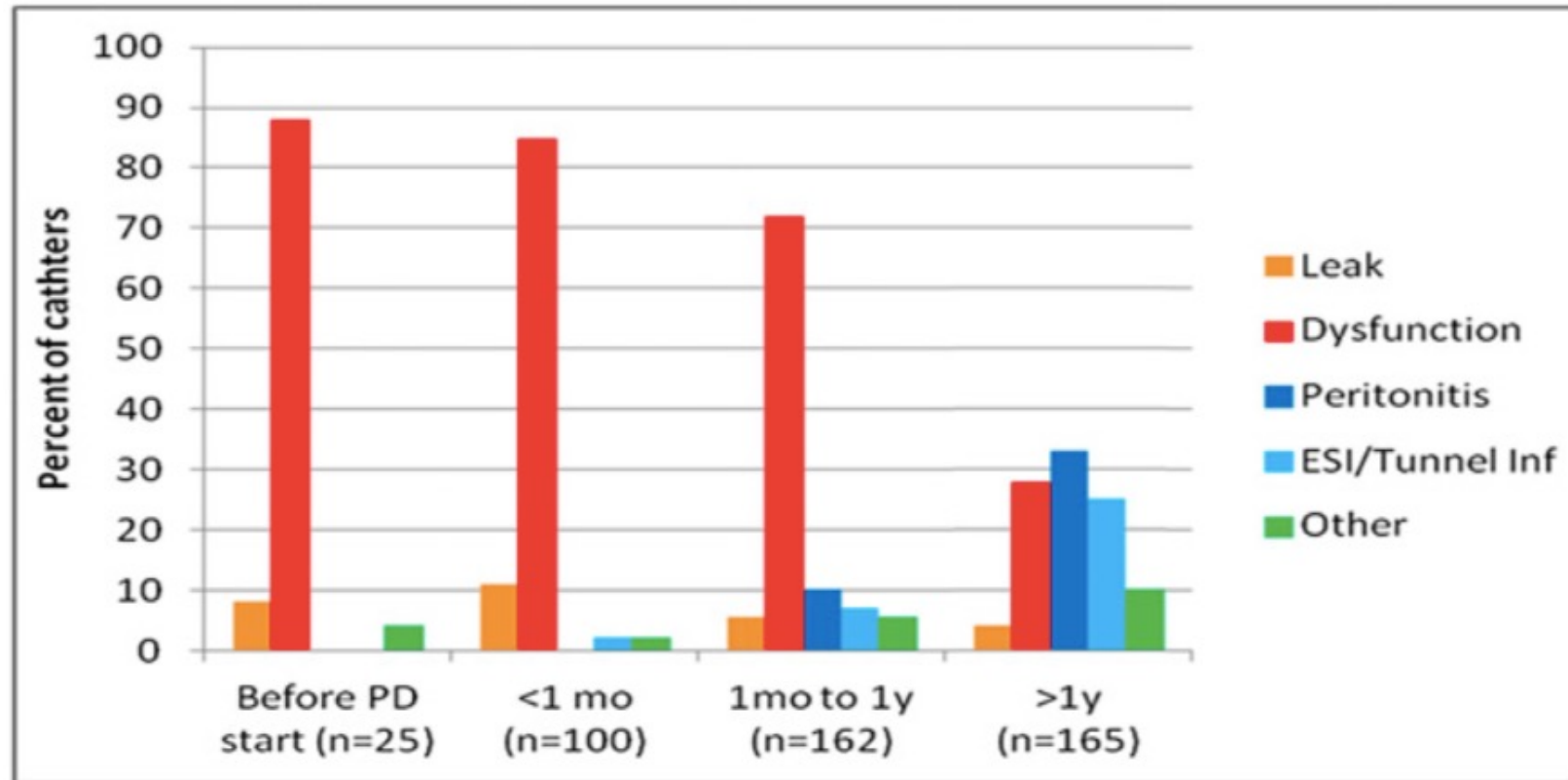
| Antimicrobial | Dose | Frequency | Max/dose |
|---|---|-------------------|---------------------------|
| Amoxicillin | 10-20 mg/kg/day | Daily | 1000 mg |
| Cephalexin | 10-20 mg/kg/day | Daily or BID | 1000 mg |
| Ciprofloxacin | 10-15 mg/kg/day | Daily | 500 mg |
| Clarithromycin | 7.5 mg/kg/day | Daily or BID | 500 mg |
| Clindamycin | 30 mg/kg/day | TID | 600 mg |
| Dicloxacillin | <40 kg 12.5-50 mg/kg/day > 40 kg 125-500 mg/dose | QID | 500 mg |
| Erythromycin (as base) | 30-50 mg/kg/day | TID or QID | 500 mg |
| Fluconazole | 1.5-3 mg/kg/day | Every 24-48 h | 200 mg |
| Levofloxacin | 10 mg/kg x 1 day, then 10 mg/kg/day | Every 48 h | Day 1 500 mg, then 250 mg |
| Linezolid | < 5 yrs 10 mg/kg/dose 5-11 yrs 10 mg/kg/dose > 12 yrs 600 mg/dose | TID BID BID | 600 mg |
| Metronidazole | 30 mg/kg/day | TID | 2000 mg/day |
| Rifampin* | 10-20 mg/kg/day | BID | 600 mg |
| Trimethoprim/Sulfamethoxazole (based on TMP) | 5-10 mg/kg/day | Daily | 80mg |

Noninfectious complications of peritoneal dialysis

Mechanical complications

- inflow/outflow problems
 - catheter malposition
 - pericatheter leak
 - hernia
-
- Children under 2 years of age or weighing less than 10 kg are at a higher risk of these complications

2453 patients - 452 PD catheter revision (2007-2015)



Mechanical catheter related problems (malfunction and leakage) doubled the risk of technique failure compared with infectious causes

Infants: More mechanical complications
Short catheter survival

| | Rinaldi ⁸ | Rahim ²⁹ | Donmez ³⁰ | Macchini ³¹ | Aksu ¹⁰ | Stringel ²² | Hooman ³⁸ | Ladd ¹⁹ | Vidal ⁷ | Phan ¹⁶ |
|--|--|--|--|---|---|--|--|---|--|--|
| Publication year | 2004, Italy | 2004, USA | 2005, Turkey | 2006, Italy | 2007, Turkey | 2008, USA | 2009, Iran | 2011, USA | 2012, Italy | 2013, USA |
| Study period | 1986-2000 | 1990-2000 | 1997-2004 | 1986-2002 | 1995-2005 | - | 1993-2006 | 1986-2008 Retrospective, single center, | 1995-2007, Registry data, | 1994-2009, Retrospective, single center |
| Number of patients | 363 (503 catheters) | 90 (127 catheters) | 53 (72 catheters) | 78 (89 catheters) | 93 (108 catheters) | 21 | 122 | 163 | 84 | 207 |
| Age | <15 years | 0-21 years | 3 days - 19 years | | 3 months-16 years | 3 months-16 years | <14 years | Mean: 6.25 ±5.58 years. | All are infants started dx <1 years | Median: 12 (range: 0-21) years |
| Insertion technique | Surgical, omentectomy in 82.4% | | Percutaneous Surgical Laparoscopic | Open surgical + omentectomy in 70% | Percutaneous | Laparoscopic + omentectomy | Surgical | All open but 1% laparoscopic, 53% partial omentectomy | Open surgical + omentectomy (97%) | Mainly open, (laparoscopic in 9%) + partial or total omentectomy in 75% |
| PD modality | | CCPD | CAPD | CAPD/CCPD | CAPD/CCPD | - | CAPD | CPD and acute (15% idiopathic acute renal failure) | CPD (70% APD 30% CAPD then APD) | CPD |
| Catheter type | Mainly double cuff staright | | Mainly double cuff swan neck-curved and straight | Mainly double cuff straight | Double cuff swan neck curved | Single cuff curled, downward or lateral exit-site | Double cuff straight or swan neck curved | Curl or staraight Quinton catheters | Mainly double cuffed, curled, downward pointing exit-site | Mostly curled |
| Timing of catheter use | | Early vs late | Early vs late | | Early vs late | After one week | Early vs. late | - | | |
| Hernia | | | 15.1% | 1.5% | No | | 20% | | | 33% for patients <1 year vs 10% for those >1 year |
| Leak | 5.8% | 14.2% | 41.5% | 2.5% | no | Several minor leaks | 15% | 13% | 3 | 18% for patients <1 year vs 3% for those >1 year |
| Kink | | | | | 7% | | | | | |
| Dislocation | 5.8% | | | 3.5% | 12% | | | 11% | 6 | |
| Malfunction (obstruction, drainage problems) | 5.3% | 21.3% | 20.8% | 5% | 7% | 7 catheter | | 36% | 9 | |
| Cuff extrusion | 4.8% | | 5.7% | | | | | | 3 | |
| Catheter exchange | 7.6% (38 catheters (17 obstruction, 14 dislocation, 4 cuff extrusion, 3 leakage) | Catheter malfunction in 11.8% of the patients, leak with infection in 1.6% | 39.6% (21 catheters from 20 patients, malfunction in 11 patients and leak in 9 patients) | 7.9% (7 catheters (6 dislocation, 1 obstruction)) | 12% (13 catheters from 11 patients (malfunction in 6, dislocation in 3, omental capture in 2, kink in 2)) | 7 interventions in 5 patients (24%) due to adhesions | Catheter obstruction in 8.7% of the patients | 63 (39%) underwent catheter revision (obstruction in 23, leak in 8, malposition in 7) | Catheter replacement in 15% of cases, all mechanical complications. 21 catheters were repositioned, due to NI complication | 46 (22%) catheters were removed for malfunction 34% adhesions, 24% leak, 17% fibrin plugs, 17% migration, and 8% other reasons |

| | Kim ^{*3} | Carpenter ¹³ | Radtke ¹² | Borzych-Duzalka ⁹ | LaPlant ¹⁴ | Radtke ¹⁵ | Imani ¹⁸ | Nikibakhsh ²⁰ |
|--|---|--|--|---|---|---|--|---|
| Publication year | 2015, Korea | 2016; USA | 2016, Germany | 2017, International | 2018, USA | 2018, Germany | 2018, USA | 2018, Iran |
| Study period | 1986-2012, Retrospective, single center | 2002-2014, Retrospective, single center | 2009-2014 Retrospective, single center | 2007-2015 Registry data International | 2005-2017, Retrospective Two centers | 2009-2015 Retrospective Two centers | Retrospective, single center, 2002-2015 | 2005-2011, Retrospective, single center |
| Number of patients | 60 patients (70 catheters) | 116 patients, 173 catheters | 60 (71 catheters) | 2453 (824 incident, 1629 prevalent) | 130 patients, 157 catheters | 122 patients, 154 catheters | 25 catheters, only < 2yrs | 56 |
| Age at dialysis initiation | 9.9±5.5 (at dx initiation) | 9.7±6.3 years (2 days to 22 years) | Median: 3.3 (0.01–15.5) years | Median 10.5 (IQR: 3.4–14.2) years | 4±5.3 years (1 day to 23 years)-46% infants | Median: 3.0 (0.01–17.1) years) | Median: 18 (7– 121) days, 60% neonate | Median: 6.5 y (1 mo – 14 years) |
| Insertion technique | - | Open (122) and laparoscopic (51) ± partial omentectomy (34%) | Open surgical | All | Mainly open and laparoscopic (n=20, 13%) + omentectomy | Open ± partial omentectomy | Laparoscopic (84%) ± omentectomy (40%) | Open surgical ± omentectomy |
| PD modality | CAPD | All CPD | 33 CPD, 37 acute | Chronic PD | Acute and CPD | 89 catheters for CPD, remaining for acute use | CPD | Acute (21) and CPD (35) (>3 weeks on PD) |
| Catheter type | Two-cuffed straight Tenckhoff, downward-pointing ES | Double cuffed catheters | One cuffed | All types | Argyle curl catheters (no straight tunnel) Upward pointing ES | One/double-cuffed curled, and straight catheters in small children (n =19), downward ES | One/double (29%)-cuffed curled (48%), and straight catheters | Swan neck coil two cuff |
| Timing of catheter use | | | | Immediately, <7 day, ≥7 day | Same day and later on --22% delayed use | | use within 3 days (48%) | Immediate use |
| Hernia | hernia (8.6%), | | | | 10% 15% in infants, and 5% in older children | | 20% hernia at catheter insertion (60% of newborns) | |
| Leak | leakage (10.0%), | | 7.1% (only in <10 kg) | 29 (%1) | 14% 21% leakage for infants vs 8% for others | 18 (11.7% of catheters) (25.5% for pts<10 kg vs 5.6% for pts>10 kg) | 32% (71% of newborns) | 5.35% |
| Dislocation | catheter tip migration (2.9%), | 7% (15% for pts < 2 y vs 5% for those >2 y | 10% | | 6% | 16 (10.4% of the catheters) | 18% | |
| Malfunction (obstruction, drainage problems) | outflow failure (14.3%), | 24% (including leak and kink) | 12.9% | 270 (%11) | 6% - adhesion | 31 (20.1% of catheters) (15 - omental trapping) | 26% | 21.4% |
| Catheter exchange | catheter malfunction, injury and oozing resulted in catheter removal in 7 (11.6%) patients – catheter exchange rate is 7.1% (n=5) | 34% of the patients had their catheter exchanged due to NI causes (dysfunction more in children < 2 years) | 17 out of 70 catheters (24.3%) needed a surgical revision within 6 months after implantation | catheter malfunction and leakage resulted in catheter exchange in 7.8% of the pts (n=192)** | 17% of the patients had their catheter exchanged (8 for leakage, 3 migration, 1 adhesion, 1 hernia) | 53 (34.4%) catheters underwent revision | 18 new catheters (72%) were inserted within 12-mo follow-up | 8.3% of CPD patients transferred to HD. NI complications are same with immediate use. |

Catheter malfunction-obstruction: 5-36% in large pediatric series (44%- in infants only)

Inflow problems – soon after placement

- intraluminal catheter occlusion (often by thrombus and fibrin)
- Catheter kinking
- Migration

Outflow failure

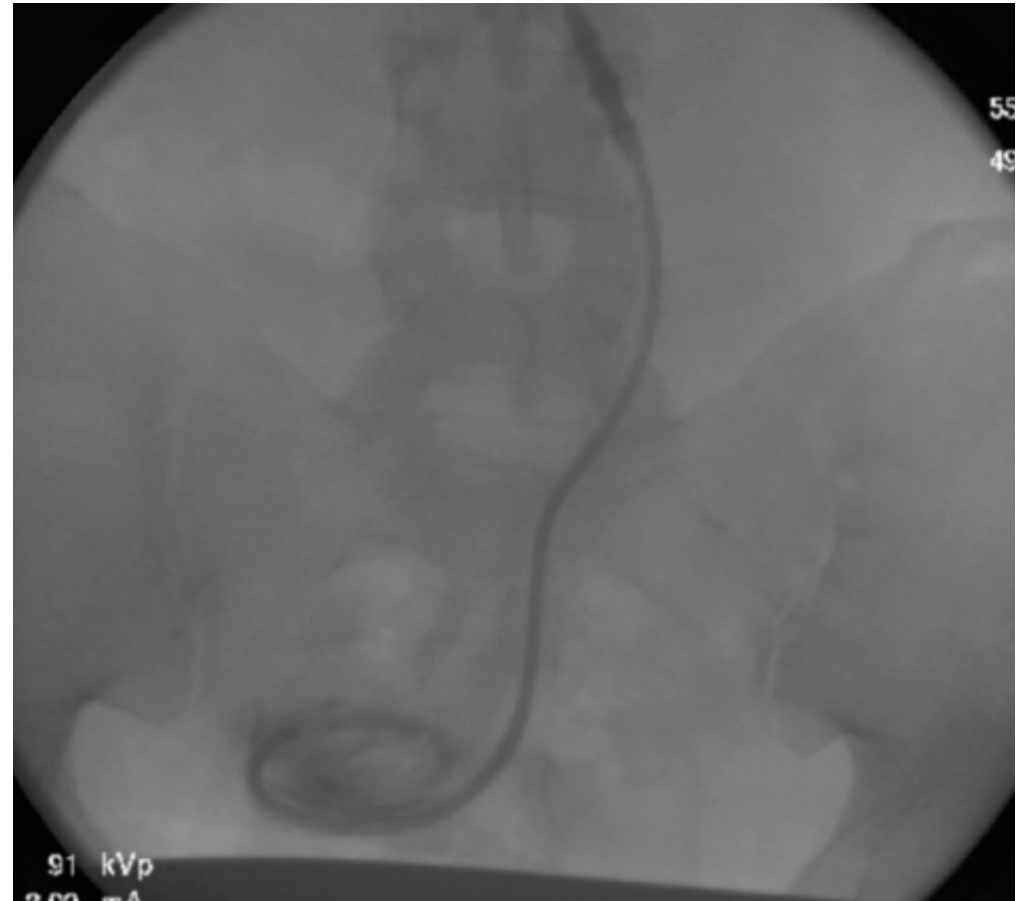
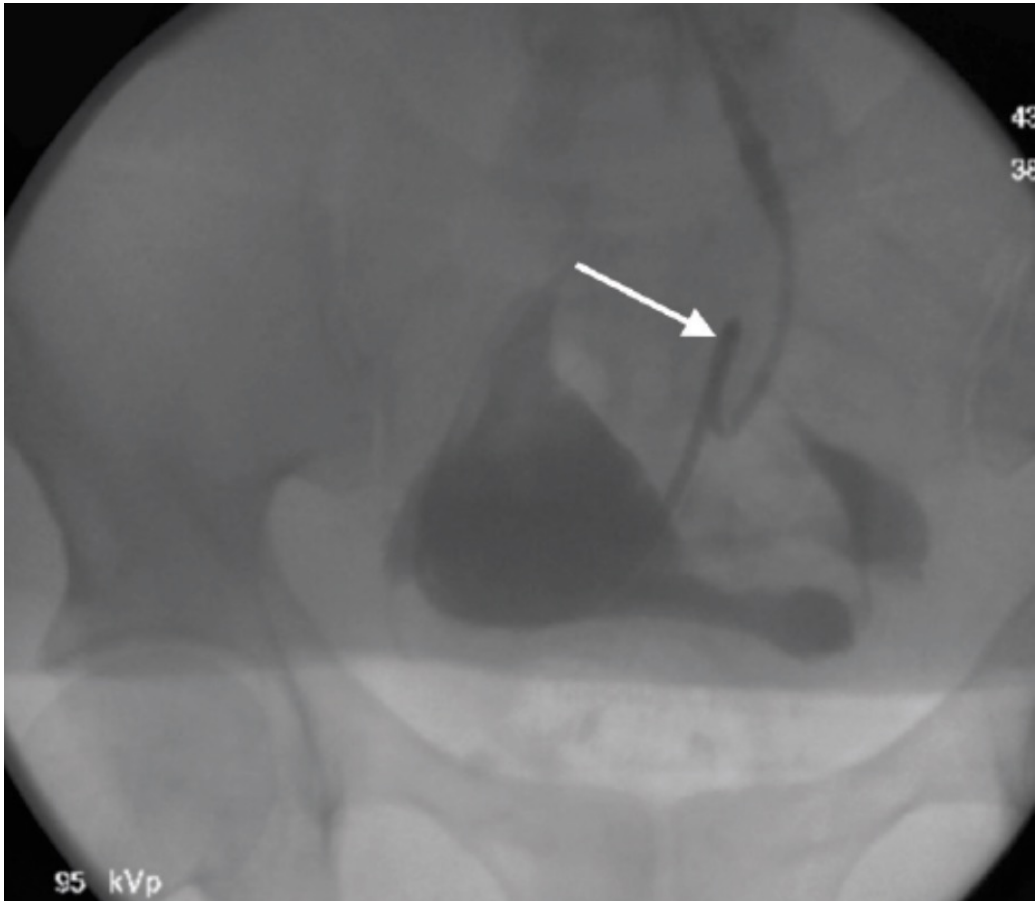
- **constipation**
- catheter malposition, tip migration
- intraluminal catheter occlusion (often by thrombus and fibrin)
- extraluminal catheter occlusion (by omentum, adhesions, epiploid fat appendices, fallopian tubes)
- catheter kinking

Prevention

- Avoid constipation by diet
- Enema, laxatives
- Saline flushing
- Cath related intervention?

Surgical expertise with the applied implantation method appears more important for preventing short and long term catheter complications than the catheter type, design or implantation technique

PD catheter kinking

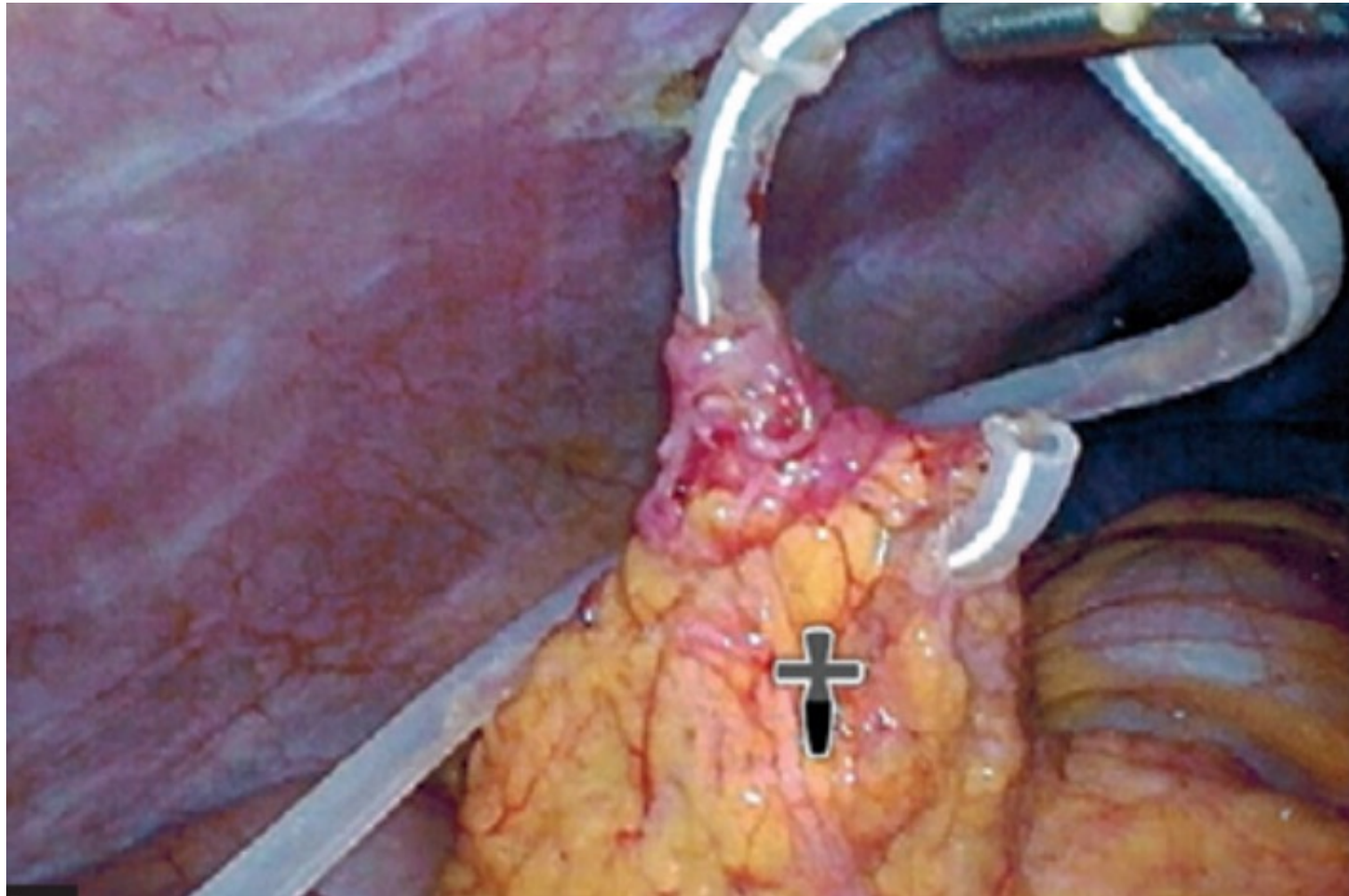


Catheter tip migration



Adult RCT - catheter fixation to the lower abdominal wall combined with a straight upward tunnel configuration and low implant position

Omental trapping



Omentectomy – Controversial data - physician's discretion in the current practice

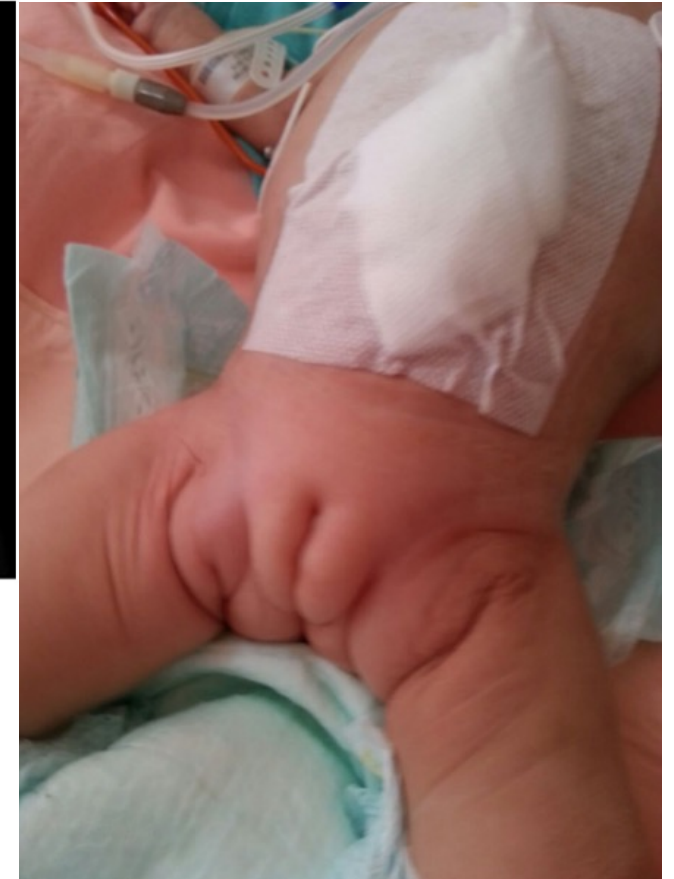
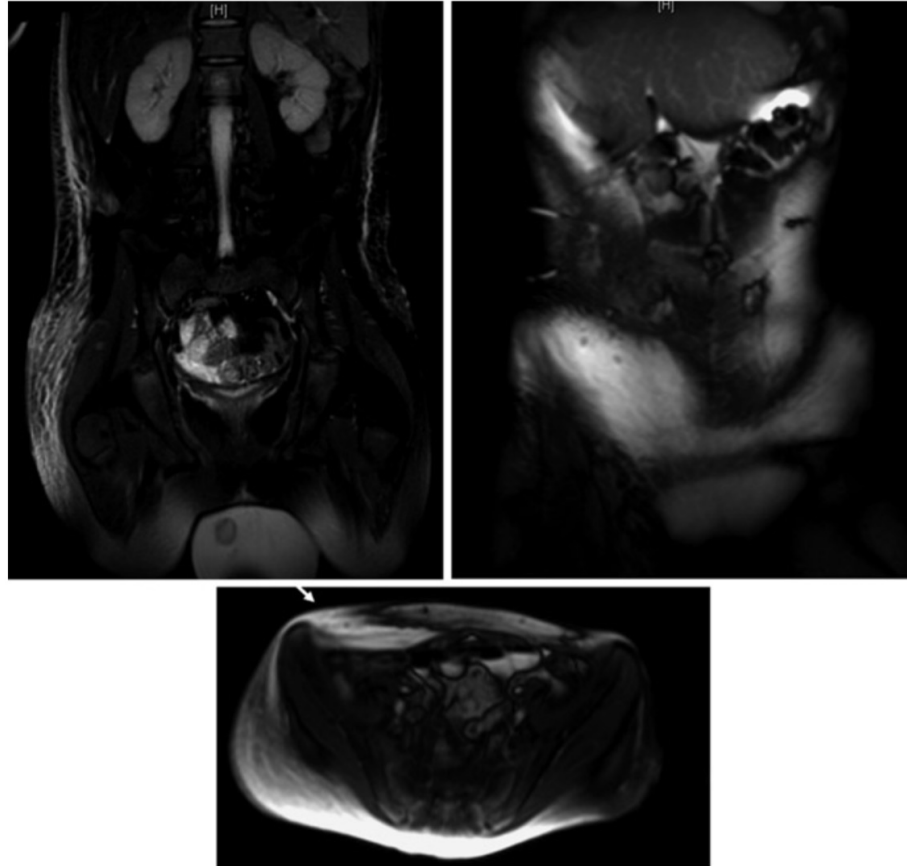
- 2-3 times reduced catheter replacement rate in patients undergoing an omentectomy (7-15% vs 23-27%)
 - LaPlant MB, (2018). Pediatr Surg Int 34:1239-1244.
 - Phan J, J Pediatr Surg 2013 48:197-202.
 - Ladd AP, et al. 2011Am J Surg 201:401-404
- Omentectomy did not change early or late mechanical complications and the re-operation rate
 - Carpenter JL, et al. J Pediatr Surg 2016; 51:730-733
 - Radtke J, et al. J Pediatr Surg 2018; 53:640-643.

Pericatheter Leakage 3-41%

Neonates, infants
Weak abdominal wall
Low subcutaneous fat
Immediate use?
Omentectomy?

Low initial dialysis volume: 300 ml/m²
Fibrin glue

Internal leakage



Hernias – up to 30%



Bakkaloglu SA, Pediatric Dialysis 2012

Bakkaloglu SA, Pediatric Dialysis 2021, courtesy of Dr. Yavaşcan

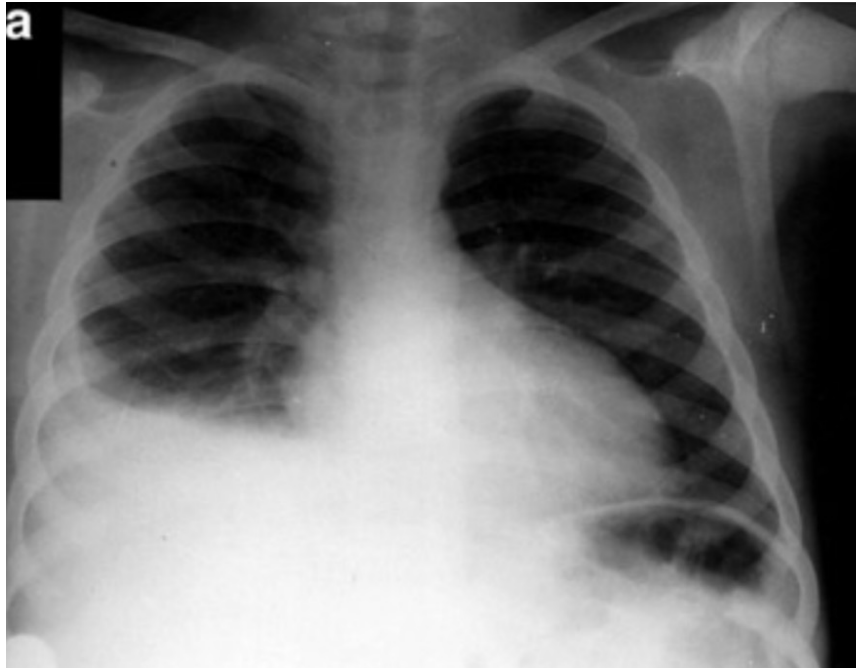
IPP
Weak sites in the abdominal wall
Small age

8-57%, newborn and small infants



Prophylactic closure of Processus vaginalis at PD cath insertion

Hydrothorax - 1.6% - 10%



Lymphatic drainage problem
Congenital diaphragmatic defect
Negative intrathoracic pressure
Positive intraabdominal pressure
IPP



Cessation of PD
Small volume PD
Thoracoscopic pleurodesis – diaphragmatic repair
Open surgery for recurrent hydrothorax cases

EPS Encapsulated peritoneal sclerosis

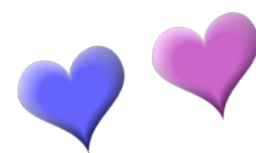


Courtesy of Dr. Ekim



Courtesy of Dr. Bakkaloglu

Long dialysis
>5 yrs of PD – incidence of EPS 6.6%
>10 yrs of PD – incidence of EPS 22%



Terminology of peritonitis

| | |
|------------------------------|---|
| Recurrent | An episode that occurs within 4 weeks of completion of therapy of a prior episode but with a different organism |
| Relapsing | An episode that occurs within 4 weeks of completion of therapy of a prior episode with the same organism or 1 sterile episode |
| Repeat | An episode that occurs more than 4 weeks after completion of therapy of a prior episode with the same organism |
| Refractory | Failure of the effluent to clear after 5 days of appropriate antibiotics |
| Catheter-related peritonitis | Peritonitis in conjunction with an exit-site or tunnel infection with the same organism or 1 site sterile |

