Peritoneal Dialysis for Acute Kidney Injury Treatment in the United States: Brought to You by the COVID-19 Pandemic

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Introduction
The coronavirus disease 2019 (COVID-19) pandemic has created unprecedented strain on health care resources in the United States. Initial reports of AKI rates from Wuhan, China ranged between 3% and 9%, although rates as high as 15% have been reported (1,2). For AKI requiring RRT, intermittent hemodialysis (HD) and continuous RRT (CRRT) have been mainstays of therapy in the United States. However, shortages in supplies, staffing, and available equipment among critically ill patients with COVID-19, particularly in the New York City area, have demanded alternative strategies such as acute peritoneal dialysis treatment for AKI (AKI-PD) that are being implemented. Here, we will review the rationale for the use of AKI-PD and describe potential advantages, criteria for patient selection, and practical considerations on the basis of initial experiences to consider when prescribing and delivering peritoneal dialysis (PD).

Rationale for Acute PD in Patients with COVID-19 and Potential Advantages
PD was routine for AKI treatment worldwide well into the 1980s. However, by the 1990s with the development of the central venous catheter for HD and the advent of CRRT, PD became rarely used in higher-resource countries to treat adult patients with AKI (3). The utility and efficacy of PD to treat patients with AKI were re-examined after the publication of a series of articles from Brazil, including a randomized trial, demonstrating that PD provided acceptable care, was not inferior to daily HD in treating acutely ill patients with AKI in terms of patient mortality, and was associated with a shorter duration of AKI and need for RRT (4,5). These findings were confirmed in a randomized trial from Saudi Arabia comparing PD with hemodialfiltration (6). AKI-PD expanded dramatically in lower-resource countries with the advent of the Saving Young Lives Program in 2012, which promoted the use of PD because of minimal infrastructural requirements, including a lack of need for water or electricity, the ease of training staff, and low costs (7–10). With the expanded PD use, the International Society for Peritoneal Dialysis (ISPD) developed guidelines for the use of PD to treat patients with AKI (3). The recommendation that PD was an acceptable form of RRT for AKI in these guidelines was supported by two meta-analyses (including a Cochrane analysis), both of which concluded that PD is not inferior to extracorporeal therapies in the management of patients with AKI (11,12).

Because PD is the original CRRT, it may be particularly suitable for hemodynamically unstable patients and for those who face challenges in establishing a reliable vascular access or with limited vascular access sites. Furthermore, unlike HD therapies, there are no concerns regarding the need for systemic anticoagulation. Emerging issues regarding a hypercoagulable state in critically ill patients with COVID-19 may pose challenges in the consistent delivery of HD or CRRT, with repeated dialysis circuit clotting making PD an attractive strategy (13). If automated acute PD prescriptions are delivered, there is a potential for reducing nursing contact with patients with COVID-19 during treatment compared with HD treatments, particularly with the use of extension tubing that allows the PD cycler and troubleshooting to take place at a distance from the patient. A recent review underscores the utility of PD to manage patients with AKI in austere environments and under conditions of duress and conflict, emphasizing the ease of implementing treatment with acceptable outcomes (14).

Concerns Regarding Acute PD and Patient Selection
At the start of PD, solute transport characteristics and ultrafiltration (UF) capacity remain unknown, which may necessitate aggressive initial empirical
prescriptions with more frequent exchanges using hyper-
tonic dialysate to maximize peritoneal UF. This is ger-
mane to patients with COVID-19 in whom greater initial fluid
removal may be often warranted (15). Unlike HD or CRRT
where fluid removal rates are visible in real time, UF often
remains unknown until the end of a PD treatment, pro-
viding an additional source of unease among critical care
team members. Maintaining enough solute clearance is an
additional concern particularly among patients who are
hypercatabolic, but using high-volume PD seems to be able
to mitigate these concerns and may be the preferable mo-
dality particularly in patients who are mechanically venti-
lated (5). Among experienced centers and using appropriate
PD catheter placement techniques (discussed below), start-
ing dwell volumes of 2.0 L have been used as suggested by
the ISPD to allow for higher earlier PD doses (3,5). Programs
with limited experience in AKI-PD or an operator new to
percutaneous PD access insertion (as may be the case with the
experience during the COVID-19 pandemic in the
United States) may first wish to have an initial test period
using lower PD dwell volumes to start.

In general, lack of knowledge and familiarity with the
performance of PD exchanges and use of automated PD by
critical care nursing staff remains a major barrier. Online or
virtual education and training support from centers with
expertise may help overcome these immediate challenges.
In particular, pediatric nurses may be an excellent resource
given the greater use of PD in children. Under extreme cases,
attending nephrologists have resorted to learning to set up
the ambulatory peritoneal dialysis (APD) cycler for their
patients with adjunct offsite nursing support. Arguably,
among staff with no acute dialysis expertise, training for
acute PD is likely more straightforward than de novo training
in HD or CRRT delivery, and consideration should be given
as such. Similar to CRRT, potassium removal is slower and
less efficient, and acute PD may not be an initial option for
patients with life-threatening hyperkalemia.

A list of potential absolute and relative contraindications
for acute PD is in Table 1. Of special note in patients with
COVID-19 is concerns of use of acute PD in patients who are
mechanically ventilated, which is not a contraindication to
acute PD. The primary concern is that increases in intra-
abdominal pressure (IAP) via PD fluid installation may
theoretically limit diaphragmatic excursion on compromise
respiratory biomechanics. Noninvasive measurement of in-
trapitoneal pressure has been described after the PD cath-
eter is in place, with typical values ranging from 10 to 16 cm
H2O. Pressures should not exceed 18 cm H2O and can be
lowered by lowering exchange dwell volumes (16,17). Yet
despite these concerns, initial reports suggest that despite
modest increases in IAP, compared with HD, PD has a min-
imal effect on respiratory biomechanics in patients who are
mechanically ventilated even at dwell volumes of 2 L
(18,19). Given that many patients with COVID-19 require
prone mechanical ventilation (although there are some
reports of successful PD in patients who are prone venti-
lated), we would suggest that an alternative dialysis mod-
dality be considered for these patients due to practical
considerations and in individuals with severe respiratory
distress where increases in IAP may potentially accelerate
the need for intubation (20,21).

In AKI-PD, the time between PD catheter placement and
initial use is short, and PD fluid leaks represent a compli-
cation seen with higher frequency compared with elective
PD starts, which usually have at least 2 weeks of healing
time from catheter insertion to first use (22,23). Neverthe-
less, the leak rates in the Brazilian and Saudi experiences
using high-volume PD therapy immediately after catheter
placement were extremely low (5,6). Other patient-related
leak risk factors include patients with diabetes, increased
body mass index, and patients on chronic immunosuppres-
sion (24). Strategies to minimize the risk of leaks include
(1) selection of initial acute PD candidates with few/no patient-
related leak risk factors, (2) optimization of PD catheter
insertion techniques to reduce the risk of leaks (discussed
below), (3) lower initial dwell volumes in the supine posi-
tion (particularly in an initial AKI-PD experience) to reduce
IAP (25), and (4) using PD as a bridge therapy from another
dialysis modality and placing the catheter early in antici-
pation of a switch to PD to allow for a longer healing period.

**PD Access Placement**

In acute PD, similar to urgent start PD, use of the catheter
within 24–48 hours demands the optimal placement tech-
nique to maximize a successful exchange, minimize the risk
of leaks, and allow for rapid escalation in dwell volumes. In
the United States, PD access is predominantly provided by
surgeons using a laparoscopic approach (26). Because of
considerations of preservation of hospital resources for the
anticipated surge of patients with COVID-19 and for the
safety of the operating room team performing laparoscopic
procedures, PD access has become difficult to arrange de-
spite Centers for Medicare and Medicaid Services designa-
tion of dialysis access procedures as essential (27,28).
Therefore, percutaneous catheter insertion with or with-
out image guidance may be considered for peritoneal access
(29,30). Although experience with percutaneous placement
is not as widespread, it is a technique that can be performed
at bedside or in the radiology suite by surgeons, interven-
tional radiologists, or interventional nephrologists who
have learned these techniques for the first time during the
COVID-19 pandemic. In addition, many surgeons have also
reverted to bedside mini-laparotomy procedures for PD
access insertion. They may be performing these procedures

| Table 1. Absolute and relative contraindications for peritoneal dialysis in AKI |
|----------------------------- |----------------------------- |
| **Factors**                      |                              |
| Recent breach of peritoneum (abdominal surgery) | Peritonitis                 |
| Bowel compromise/inflammation  | Severe hyperkalemia          |
| Severe respiratory failure and pulmonary edema | Shock liver and/or severe lactic acidosis\* |
| Ascites and high intra-abdominal pressure | Prone ventilation |

\*Only a relative contraindication with lactate-buffered (not bicarbonate-buffered) peritoneal dialysis solutions. Bicarbonate-buffered solutions are not currently available in the United States.
using this method for the first time who would have traditionally exclusively placed PD catheters via a laparoscopic technique. During access placement, leak risk is minimized with (1) the use of a purse-string suture to secure the deep cuff, which should be placed in the rectus muscle (31). (2) A paramedian over a midline incision into the peritoneal cavity may further reduce leak risk by providing better adherence of the deep cuff to the lateral rectus muscle laterally compared with the thinner medial tissues of the linea alba, although this remains controversial (32). For all procedures, prophylactic antibiotics at the time of PD access insertion should be used in keeping with ISPD guidelines to reduce early peritonitis risk (33). Local expertise and operator experience with the technique being considered should be the main drivers for the method of PD access insertion. One of the main advantages for acute PD is that the PD catheter may also serve as a long-term access should the patient fail to recover from the AKI episode.

Acute PD Protocol and Prescription Considerations

An acute PD prescription must carefully balance the metabolic and UF needs of the patient while minimizing the risk of treatment-related complications. What dose to be delivered is controversial and has been poorly studied. The Brazilians targeted daily Kt/V of approximately 0.6/d, which may be necessary in very catabolic patients, but the ISPD has suggested that daily Kt/V urea of 0.3 may be adequate for many patients with AKI (3,5). A dose estimation guide is provided in Tables 2–4, although Kt/V urea in PD may not be the appropriate metric for the dose of dialysis in AKI. Furthermore, consideration must be given to local resources available, including nursing capability, familiarity with both manual and automated PD, cycle supply, dialysate supply, and nephrologists’ ability to identify and manage complications.

A sample PD prescription and protocol are provided in Figure 1. Bowel hygiene is important to optimize catheter function, with a bowel routine protocol in place from the time of placement and over the course of therapy. Both automated and manual PD exchanges are possible with an acute PD prescription in the supine position to minimize the risk of increased IAP and leaks. If manual exchanges are performed, continuous APD systems can be used with standard equipment or using the manifold and clamps to minimize the number of connections and disconnections needed. Automated PD prescriptions need not be necessarily prescribed for overnight treatments alone; they may be set up for continuous (24-hour) treatments and have been largely used for bed-bound patients and patients who are mechanically ventilated. In these patients, the PD cycler can be set up for one 24-hour treatment and 60- to 240-minute exchanges used as clinically indicated. With excessively short APD dwell times and hypertonic dialysate, there is a greater risk of sodium sieving, particularly with hypertonic solutions leading to excessive free water loss (in the absence of sodium removal), and biochemistry should be reviewed for rises in serum sodium (3,34). With frequent automated PD cycling, hypokalemia may also ensue, necessitating intraperitoneal and/or intravenous potassium supplementation. More frequent cycling may also promote a greater risk of APD alarms overnight, and as a result, less frequent cycles and using tidal PD may be advantageous overnight with fewer/no staff available to troubleshoot these alarms. If a leak develops, temporary cessation of PD may be needed and has been introduced as early as within 24 hours of rest using lower dwell volumes. If persistent, catheter replacement may be necessary using the techniques described above to reduce the risk of leaks. Drug dosing in AKI-PD has not been well established in particular for antimicrobials and could be potentially extrapolated from the CRRT literature. Where possible, antibiotic drug levels should be measured and followed.

For all PD exchanges, intraperitoneal heparin supplementation (500–1000 μl) has been given either prophylactically to prevent intraperitoneal fibrin formation or as needed on

**Table 2. Peritoneal dialysis treatment for AKI dialysis orders: dialysis prescription—automated peritoneal dialysis order (first 24-hour prescription)**

| Parameter                     | Weight (≤70 kg) | Weight (>70 kg) |
|-------------------------------|----------------|-----------------|
| Fill volume, ml               | 1000           | 1500            |
| Time, h                       | 8–24           | 8–24            |
| No. of cycles                 | 8–24           | 8–24            |
| Total therapy volume, ml      | 8000–24,000    | 12,000–36,000   |
| Dwell time per exchange, h    | 1              | 1               |

For intensive care unit, 16–24 hours. For patients on floor, start with 8–12 hours.

**Table 3. Peritoneal dialysis treatment for AKI dialysis orders: prescription—after 24–48 hours (no leaks), increase dwell volume and time**

| Parameter                     | Weight (≤70 kg) | Weight (>70 kg) |
|-------------------------------|----------------|-----------------|
| Fill volume, ml               | 1500           | 2000            |
| Time, h                       | 8–24           | 8–24            |
| No. of cycles                 | 4–12           | 4–12            |
| Total therapy volume, ml      | 6000–18,000    | 8000–24,000     |
| Dwell time per exchange, h    | 2              | 2               |

For continuous automated peritoneal dialysis, consider 2-hour dwell time per exchange.

**Table 4. Peritoneal dialysis treatment for AKI dialysis orders: dextrose concentration (is on the basis of volume status and ultrafiltration requirement)**

| Dextrose Concentration     | No. of Liters |
|----------------------------|---------------|
| 1.5% (if no fluid overload)|               |
| 2.5% (if mild or moderate fluid overload) |               |
| 4.25% (if severe fluid overload) | 4.25% solution can remove up to 1 L of fluid in 4 h |
the basis of appearance of effluent fibrin to maintain PD catheter patency. Heparin is too large to cross the peritoneal membrane, and therefore, it is not contraindicated in patients with bleeding diatheses but is contraindicated in patients with heparin-induced thrombocytopenia, where intraperitoneal heparin has been reported to elicit an immunologic response (35). With sluggish PD catheter function, drain pain, prolonged inflow or outflow times, or excessive automated PD cyclers low drain alarms, tidal PD (leaving a fixed residual volume of dialysis solution) during each exchange may be required. Spent dialysis fluid can be discarded with the same precautions as used for other bodily fluids (i.e., urine) among patients who are COVID-19 positive, although viral replication of COVID-19 has been recently identified in PD effluent (36).

PD for AKI is an established RRT with acceptable outcomes in pediatric patients with AKI and in adult patients outside of the United States (37). It is our hope that the renewed interest in the treatment of PD for AKI in adult patients in the United States during the COVID-19 pandemic is accompanied by increased proficiency and comfort with providing and offering this treatment modality and encouraging initial reports. For programs with established expertise in managing patients on maintenance PD or with expertise in using PD for urgent starts in the late-referred patient with ESKD, PD for AKI may be less of a leap compared with programs with little experience in maintenance PD where such an endeavor may be more challenging and perhaps ill advised. In such patients or where there is reluctance among the critical care team, use of urgent start PD in the late-referred patients with ESKD or greater use of PD in sub-AKI or as a bridging therapy from HD may offload HD and CRRT resources reserved for critically ill patients. For an AKI-PD program to be successful, it will require a team approach centered around support from the critical care team, tenets of PD access insertion reliability and speed, nursing expertise, standardization and implementation of protocols, and evidence-based practice (where available). Initial candidates may want to be considered carefully and more restrictively, particularly in initially choosing lower acuity candidates and from a PD access perspective, candidates with no prior major abdominal surgery or scarring. If PD is not meeting the patient’s goals for RRT for AKI after two treatments, it is important to swiftly consider an alternate dialysis modality. As patients are ready for discharge from the hospital with ongoing AKI requiring RRT, discharge planners will need to work with outpatient dialysis facilities to transition the patient to outpatient PD. Currently, in the United States, few insurance providers pay for AKI-PD; therefore, the patient management team will need to be involved so that coverage can be guaranteed prior to discharge via the health plan or through an agreement between the hospital and the dialysis provider. This safe transition should also include an in-home assessment to ensure the patient’s long-term success on the modality after discharge.

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Author Contributions
F.O. Finkelstein, M. Naljayan, and J. Perl conceptualized the study; J. Perl was responsible for resources; and V. Aggarwal, J.H. Crabtree, F.O. Finkelstein, M. Naljayan, J. Perl, and V. Srivatana wrote the original draft and reviewed and edited the manuscript.

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References
1. Huang C, Wang Y, Li X, Ren L, Zhao J, Hu Y, Zhang L, Fan G, Xu J, Gu X, Cheng Z, Yu T, Xia J, Wei Y, Wu W, Xie X, Yin W, Li H, Liu M, Xiao Y, Gao H, Guo L, Xie J, Wang G, Jiang R, Gao Z, Jin Q, Wang J, Cao B: Clinical features of patients infected with 2019 novel coronavirus in Wuhan, China [published correction appears in Lancet 395: 496, 2020]. Lancet 395: 497–506, 2020
2. Naciker S, Yang CW, Hwang SJ, Liu BC, Chen JH, Jha V: The novel coronavirus 2019 epidemic and kidneys. Kidney Int 97: 824–828, 2020
3. Cullis B, Abdelraheem M, Abrahams G, Balbi A, Cruz DN, Frishberg Y, Koch V, McCulloch M, Munangulu A, Nourse P, Pequito-Filho R, Ponce D, Wardby B, Yeates K, Finkelstein FO: Peritoneal dialysis for acute kidney injury. Perit Dial Int 34: 494–517, 2014
4. Ponce D, Berbel MN, Regina de Goes C, Almeida CT, Balbi AL: High-volume peritoneal dialysis in acute kidney injury: Indications and limitations. Clin J Am Soc Nephrol 7: 887–894, 2012
5. Gabriel DP, Caramori JT, Martim LC, Barretti P, Balbi AL: High-volume peritoneal dialysis vs daily hemodialysis: A randomized, controlled trial in patients with acute kidney injury. Kidney Int Suppl 73(Suppl): S87–S93, 2008
6. Al-Hwisesh A, Abdul-Rahman I, Finkelstein F, Divino-Filho J, Qutub H, Al-Audah N, Abdelrahman A, El-Fakhirny N, Nasr Eldin M, El-Salamony T, Noor A, Al-Shahrani M, Al-Otaibi K: Acute kidney injury in critically ill patients: A prospective randomized study of total peritoneal dialysis versus continuous renal replacement therapy. Ther Apher Dial 22: 371–379, 2018
7. Smoyer WE, Finkelstein FO, McCulloch M, Carter M, Brussel- mans A, Feehally J: “Saving Young Lives” with acute kidney injury: The challenge of acute dialysis in low-resource settings. Kidney Int 89: 254–256, 2016
8. Smoyer WE, Finkelstein FO, McCulloch M, Carter M, Brussel- mans A, Feehally J: Saving Young Lives: Provision of acute dialysis in low-resource settings. Lancet 386: 2056, 2015
9. Finkelstein FO, Smoyer WE, Carter M, Brusselmann A, Feehally J: Peritoneal dialysis, acute kidney injury, and the Saving Young Lives program. Perit Dial Int 34: 478–480, 2014
10. Abdou N, Antwi S, Kofi LA, Lalaya F, Adabayeri VM, Nyah N, Palmer D, Brusselmann A, Cullis B, Feehally J, McCulloch M, Smoyer W, Finkelstein FO: Peritoneal dialysis to treat patients with acute kidney injury—the Saving Young Lives experience in West Africa: Proceedings of the Saving Young Lives session at the First International Conference of Dialysis in West Africa, Dakar, Senegal, December 2015. Perit Dial Int 37: 155–158, 2017
11. Liu L, Zhang L, Liu GJ, Fu P: Peritoneal dialysis for acute kidney injury. Cochrane Database Syst Rev 12: CD011457, 2017
12. Chionh CY, Soni SS, Finkelstein FO, Ronco C, Cruz DN: Use of peritoneal dialysis in AKI: A systematic review. Clin J Am Soc Nephrol 8: 1649–1660, 2013
13. Tang N, Li D, Wang X, Sun Z: Abnormal coagulation parameters are associated with poor prognosis in patients with novel coronavirus pneumonia. J Thromb Haemost 18: 844–847, 2020
14. Gorbatkin C, Bass J, Finkelstein FO, Gorbatkin SM: Peritoneal dialysis in austere environments: An emergent approach to renal failure management. West J Emerg Med 19: 548–556, 2018
15. Phua J, Weng L, Ling L, Egi M, Lim CM, Divatia JV, Shrestha BR, Arabi YM, Ng J, Gomersall CD, Nishimura M, Koh Y, Du B: Asian Critical Care Clinical Trials Group: Intensive care management of coronavirus disease 2019 (COVID-19): Challenges and recommendations [published online ahead of print April 6, 2020]. Lancet Respir Med doi:10.1016/S2213-2600(20)30161-2
16. Mathieu B, Pillet A; Alir Nursing Team; Measurement of hydrosystatic intraperitoneal pressure. Adv Perit Dial 10: 59–62, 1994
17. Durand PY: Measurement of intraperitoneal pressure in PD patients. Perit Dial Int 25: 333–337, 2005
18. Almeida CP, Ponce D, de Marchi AC, Balbi AL: Effect of peritoneal dialysis on respiratory mechanics in acute kidney injury patients. Perit Dial Int 34: 544–549, 2014
19. Almeida CP, Balbi AL, Ponce D: Effect of peritoneal dialysis vs. haemodialysis on respiratory mechanics in acute kidney injury patients. Clin Exp Nephrol 22: 1420–1426, 2018
20. Kilsnick A, Souweine B, Fileitre M, Wauquier JP, Gazzu NY, Detex P, Baguet JC: Peritoneal dialysis in a patient receiving mechanical ventilation in prone position. Perit Dial Int 18: 536–538, 1998
21. Meng L, Qiu H, Wan L, Ai Y, Xue Z, Guo Q, Deshpande R, Zhang L, Meng L, Tong C, Liu H, Mu D: Ventilation and ventilation amid the COVID-19 outbreak: Wuhan’s experience [published online ahead of print March 26, 2020]. Anesthesiology doi:10.1097/ALN.0000000000003296
22. Ghafari A: Urgent-start peritoneal dialysis: A quality improvement report. Am J Kidney Dis 59: 400–408, 2012
23. Iversen F, Povlsen JV: Can peritoneal dialysis be applied for unplanned initiation of chronic dialysis? Nephrol Dial Transplant 29: 2201–2206, 2014
24. Del Peso G, Bajo MA, Costero O, Hevia C, Gil F, Díaz C, Aguilera A, Selgas R: Risk factors for abdominal wall complications in peritoneal dialysis patients. Perit Dial Int 23: 249–254, 2003
25. Tvardowskij ZV, Prowant BF, Nolph KD, Martinez AJ, Lampton LM: High volume, low frequency ambulatory ambulatory peritoneal dialysis. Kidney Int 23: 64–70, 1983
26. Wilkie M, Zhao J, Biber B, Bowes E, Crabtree J, Fluck R, Fukasawa M, Jain A, Pisoni R, Quinn R, Teitelbaum I, Perl J; Working Group on behalf of PDOPPS Catheter Access and Function W: International variation in peritoneal dialysis (PD) catheter practices: Preliminary results from the Peritoneal Dialysis Outcomes and Practice Patterns Study (PDOPPS). Nephrol Dial Transplant 32: iii297–iii299, 2017
27. SAGES: SAGES and EAES recommendations regarding surgical response to COVID-19 crisis, 2020. Available at: https://www.sages.org/recommendations-surgical-response-covid-19/. Accessed March 25, 2020
28. White D: Critical clarification from CMS: PD catheter and vascular access placement is essential. Kidney News Online, 2020. Available at: https://www.kidneynews.org/policy-advocacy/leading-edge-critical-clarification-from-cms-pd-catheter-and-vascular-access-placement-is-essential. Accessed March 25, 2020
29. Abdel-Aal AK, Dybbro P, Hathaway P, Guest S, Neuwirth M, Krishnamurthy V: Best practices consensus protocol for peritoneal dialysis catheter placement by interventional radiologists. Perit Dial Int 34: 481–493, 2014
30. Al-Hwisesh A: Percutaneous peritoneal dialysis catheter insertion by a nephrologist: A new, simple, and safe technique. Perit Dial Int 34: 204–211, 2014
31. Stegmayr BG: Three purse-string sutures allow immediate start of peritoneal dialysis with a low incidence of leakage. Semin Dial 16: 346–348, 2003
32. Eijersen E, Steven K, Lokkegaard H: Paramedian versus midline incision for the insertion of permanent peritoneal dialysis catheters. A randomized clinical trial. Scand J Urol Nephrol 24: 151–154, 1990
33. Li PK, Szeto CC, Piraino B, de Arteaga J, Fan S, Figueiredo AE, Fish DN, Goffin E, Kim YL, Salzer W, Struijk DG, Teitelbaum I, Johnson DW: ISPD peritonitis recommendations: 2016 Update on prevention and treatment. *Perit Dial Int* 36: 481–508, 2016

34. Gomes AM, Fontán MP, Rodríguez-Carmona A, Sastre A, Cambre HD, Muñiz AL, Falcón TG: Categorization of sodium sieving by 2.27% and 3.86% peritoneal equilibration tests—a comparative analysis in the clinical setting. *Nephrol Dial Transplant* 24: 3513–3520, 2009

35. Kaplan GG, Manns B, McLaughlin K: Heparin induced thrombocytopaenia secondary to intraperitoneal heparin exposure. *Nephrol Dial Transplant* 20: 2561–2562, 2005

36. Nouvier M, Chalencon E, Novel-Catin E, Pelletie S, Hallonet P, Charre C, Koppe L, Fouque D: First viral replication of Covid-19 identified in the peritoneal dialysis fluid of a symptomatic patient. *Bulletin de la Dialyse a Domicile* 3: 49–50, 2020

37. Vasudevan A, Phadke K, Yap HK: Peritoneal dialysis for the management of pediatric patients with acute kidney injury. *Pediatr Nephrol* 32: 1145–1156, 2017