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CASE REPORT

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INTENSIVE CARE

Management of peritoneal dialysis in an adolescent with septic shock, AKI, multi-organ failure and COVID-19: a case report

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ABSTRACT

The kidneys is the most commonly injured organs in sepsis, which might manifest as acute kidney injury (AKI). Peritoneal dialysis (PD) is the preferred modality of renal replacement therapy (RRT), especially in the developing countries. Furthermore, the COVID-19 pandemic has made hemodialysis (HD) unavailable due to infection concerns. We present a critically-ill 16-year-old male with morbid obesity (BMI 43.3 kg/m²), diabetic ketoacidosis, acute pancreatitis, acute kidney injury, and COVID-19 pneumonia. The patient looked somnolent on admission with very high random blood glucose and hypernatremia. Blood gas analysis showed metabolic acidosis. The patient was treated with 1 g/day Meropenem, 0.5 μ g/kg/min norepinephrine, 0.04 unit/kg/h vasopressin, and correction of electrolyte imbalance and metabolic acidosis. After 48 h, the diuresis and the creatinine levels worsened. We placed a Tenckhoff catheter and started PD. The patient showed improvement after 72 h of peritoneal dialysis.

This case report highlighted the use of PD as a modality of RRT in sepsis-induced AKI, multi-organ failure, and COVID-19. Although it remains controversial, we observed an improvement in diuresis and creatinine levels following PD. PD is more cost-effective and provides a similar outcome compared to other modalities.

Abbreviations: AKI - Acute Kidney Injury; BMI- Body Mass Index; CRRT- Continous Renal Replacement Therapu; GCS-Glasgow Coma Scale; HD- Hemodialysis; ICU- Intensive Care Unit; PD- Peritoenal Dialysis; RRT- Renal Replacement Therapy; S-AKI- Sepsis-Associated Acute Kidney Injury; SIRS- Systemic Inflammatory Response Syndrome

Key words: Septic shock; Acute kidney injury; Peritoneal dialysis

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1. INTRODUCTION

Sepsis is the leading cause of acute kidney injury (AKI). The most frequent source of infection that causes sepsis is the respiratory tract, including COVID-19 pneumonia.¹ Kidney hypoperfusion, hemodynamic instability, inflammatory mediators, and kidney vascular dysfunction and inflammation result in acute tubular necrosis, ischemia-reperfusion damage to the glomeruli, inflammation of the nephrons, oxidative stress, renal tubular injury, and apoptosis of the tubular and renal mesenchyme. These conditions are known to cause sepsis-induced AKI.^{2–4} Lack of perfusion manifests as lactic acidosis, oliguria, or alteration in the mental status.¹



Figure 1: Chest X-ray shows increased bronchovascular markings and opacities in the lung fields.

The treatment of AKI is renal replacement therapy (RRT), with either continuous RRT, hemodialysis (HD), or peritoneal dialysis (PD), with the latter being an option for kidney failure. The geographic location, socio-economic conditions, and clinical manifestations of the patient appear to affect the selection of the modality.⁵ PD is sometimes the primary dialysis choice in low- to middle-income countries. It can remove waste products from the body using dialysate and the peritoneum as a filter for this process.⁴ This case report highlights the use of PD as a treatment modality for sepsis-associated acute kidney injury (S-AKI) in an adolescent with multi-organ failure. The objective is to remind the physicians treating sepsis-induced renal failure about the usefulness of PD when other options are restricted or non-available.

2. CASE REPORT

A 16-year-old male presented to the emergency department with sudden loss of consciousness. Glasgow Coma Scale (GCS) was 8 for the last 5 h prior to hospitalization, and this SARS CoV-2 test was positive. His body temperature was 39.8°C, heart rate 144 beats/min, respiratory rate 30 breaths/min, blood pressure 83/52 mmHg, and oxygen saturation at 97% with 4 L/min oxygen administered via a nasal cannula. Physical examination revealed him to be obese with BMI 43.3 kg/m², and rales at the basal areas of the lungs. Laboratory results were remarkable for undetectable very high blood glucose level, hypernatremia (serum sodium 156 mmol/L), hyperchloremia (120 mEq/L), and elevated lactate level (2.8 mmol/L). His blood gas analysis revealed pH 6.9, PCO₂ 23.4 mmHg, PaO₂ 122 mmHg, HCO3⁻ 10 meq/L, an elevated creatinine 2.3 mg/dl and procalcitonin 12 ng/ml. Amylase was 566 U/L, and lipase was 128 U/L. The patient presented with hypotension unresponsive to fluid resuscitation. The patient was intubated and ventilated, a chest X-ray obtained (Figure 1) and transferred to the intensive care unit (ICU).

The patient was diagnosed with septic shock and sepsis induced AKI based on the clinical and laboratory findings. The patient's condition was accompanied by morbid COVID-19 pneumonia, diabetic ketoacidosis, acute pancreatitis, and obesity.

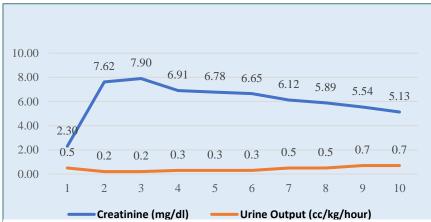
The ventilator setting were: AC/PC mode with FiO₂ 35%, Pins 12 cmH₂O, PEEP 5 cmH₂O, RR 15/min, and PS 12 cmH₂O. Norepinephrine infusion was started to maintain adequate blood pressure, followed by vasopressin infusion. After normalizing the acidemia and electrolyte imbalances, vasopressors could be tapered off. The patient was started on meropenem, ciprofloxacin, and remdesivir while sedated with ketamine 40 µg/kg/h. By the second day of hospitalization, a net positive fluid balance of 2.5 L was noted, and the hypernatremia persisted (serum sodium of 153 mmol/L). We started furosemide infusion 5 mg/h. After 3 h of furosemide, the patient's urine output remained at only 0.2 ml/kg, and the creatinine level rose to 7.62 mg/dl. We placed Tenckhoff catheter in the rectovesical pouch and PD was started for the next 72 h, with dialysate solution containing insulin, potassium, heparin, and ceftazidime.

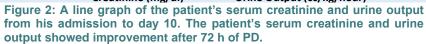
After 72 h of PD, the patient showed an improvement in diuresis of 0.7 ml/kg/h. His creatinine level dropped to 6.12 mg/dl and 5.13 mg/dl the subsequent day. He remained hospitalized in the ICU for nine days.

3. DISCUSSION

The sepsis-induced AKI in our patient may have multifactorial causes, such as COVID-19 pneumonia, diabetic ketoacidosis, acute pancreatitis, and obesity. Diabetic ketoacidosis is associated with acute pancreatitis, which can lead to beta-cell dysfunction and transient insulin deficiency and increase insulin resistance due to systemic inflammation. Patients with diabetic ketoacidosis are at greater risk of developing systemic inflammatory response syndrome (SIRS).⁶

he patient was obese class-III, with a BMI of 43.3 kg/m². Research conducted by Ganda et al. found that the prevalence of S-AKI patients with malnutrition was higher than patients with normal nutritional states. Malnutrition affects systemic function, including immunity, cell damage, and membrane permeability of the intestinal mucosa, which promotes bacterial translocation. This condition increases the risk of pneumonia, sepsis, and other infections, resulting in





higher mortality, increased length of hospitalization and care $cost.^1$

RRT modalities for pediatric AKI have evolved from PD, HD, continous renal replacement therapy (CRRT), to sustained low-efficiency dialysis (SLED).⁷ The socioeconomic and clinical characteristics of the patient are factors that influence the choice of treatment with PD. Several retrospective cohort studies have shown that PD can be conducted in pediatric patients with hemodynamic instability and multi-organ failure requiring vasopressors,⁵ as in our patient and the PD therapy we used.

We used a dialysate with a combination of 2 L of 2.5% dextrose with 1000 U heparin, 10 mEq potassium, 10 U Novorapid[®], and 250 mg ceftazidime. In PD, the peritoneum functions as a filter and the dialysate solution

will remain in the peritoneum for 40 min to one hour and be exchanged every 6 h. The procedure of PD is technically straightforward, does not require vascular access or expensive equipment, and is more practical and safer an RRT.8 as Additionally, it can be performed with or without a machine. + PD peel-away Tenckhoff with catheters have a substantially reduced risk of complication than CRRT.4,9

A prospective study showed that early PD therapy in patients with S-AKI was correlated with early renal improvement while also

being associated with decreased PD duration, early dialysis termination, and fewer procedural risks.⁴ PD does not require extracorporeal circulation and does not involve blood exposure to synthetic membranes, so the hemodynamic tolerance is relatively good. These factors potentially contribute to the early recovery of kidney function.¹⁰ Our patient showed an improvement in diuresis of 0.7 ml/kg/h, and a fall in creatinine level to 5.5 mg/dl.

Prior to the COVID-19 pandemic, CRRT was the first choice for RRT in S-AKI, especially in hemodynamically unstable patients. However, after the pandemic, numerous hospitals found it difficult to allocate HD machines for many COVID-19 with AKI due to cross-infection risks.⁹

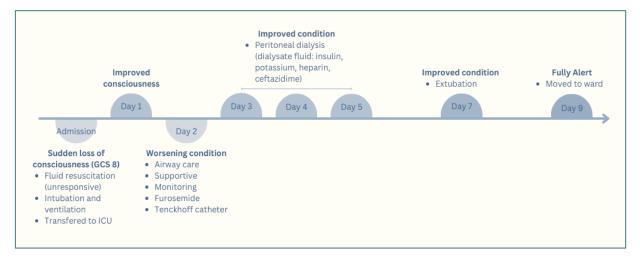


Figure 3: Patient's condition timeline. The patient came with a sudden loss of consciousness and presented unresponsive hypotension to fluid resuscitation, then the patient was moved to ICU. The peritoneal dialysis was started on day 3 for the next 72 h,

Some concerns regarding PD have been raised, including whether appropriate clearance and ultrafiltration can be accomplished, and the peritonitis risk and the impact of protein loss and glucose absorption. These factors lead to PD being practiced only in developing countries where advanced therapies are still minimal. However, recent data showed that mortality and complication rates of PD in AKI have decreased even further and there is no reason to believe that other modalities offer any advantage over PD in regards to the outcome.¹⁰

4. CONCLUSION

Peritoneal dialysis can be one of the treatment options for AKI, especially in patients with sepsis, in countries where advanced therapies are not readily available. It is cost-effective, can easily be performed, and does not require vascular access and expensive equipment. It offers outcomes comparable to hemodialysis. This therapy needs to be evaluated further to weigh its risks and outcomes.

5. Ethical committee approval

Ethical approval is not required for this study in accordance with local or national guidelines.

6. Consent to participate

Written informed consent for publication was obtained from the patient's biological father for publication of this case report with any accompanying images.

7. Availability of Data and Materials

Numerical data is available from the authors upon reasonable request and with permission of the hospitsl administration.

8. Conflict Interests

The authors declare that they have no conflicting interests.

9. Acknowledgments

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10. Author's Contributions

Conceptualization: RS. Writing-original draft: DSA. Writingreview & editing: DSA, JJ. All authors read and approved the final manuscript.

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