

A 10-Year Journey of Continuous Ambulatory Peritoneal Dialysis (CAPD) Peritonitis: Lessons from Dr. Saiful Anwar Hospital, Malang, Indonesia



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Disclosure of Conflict of Interest

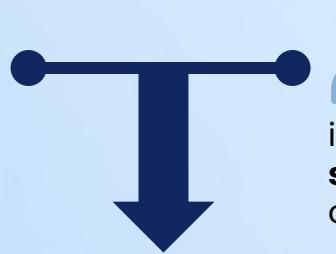
All authors have nothing to declare for this study

Background



Continuous Ambulatory Peritoneal Dialysis (CAPD)

is an effective renal replacement therapy for patients with **end-stage renal disease (ESRD)**



Peritonitis

is defined as **inflammation of the serous membrane** lining the abdominal cavity and its organs

Dialysis-related peritonitis remains a major complication and a key challenge to the long-term success of peritoneal dialysis

- **Mortality Rate 5%**

Mortality in 16% of infected patients

- Severe, refractory, or prolonged CAPD peritonitis → **Peritoneal dialysis failure, switching to hemodialysis, and increased mortality**

OBJECTIVE

To analyze the etiological characteristics of CAPD peritonitis, a major cause of mortality over ten years at Dr. Saiful Anwar Hospital, Malang

1. Bach N, et al. *BMC Nephrol*. 2025;26(1).
2. Nardelli L, et al. *BMC Nephrol*. 2024;25(1):1-9.
3. Song P, et al. *Front Med (Lausanne)*. 2022;9:799110.

Methods

Study Design: Retrospective descriptive study

Study Site: CAPD Unit, Dr. Saiful Anwar General Hospital, Malang, Indonesia.

Data Source: Medical record review of CAPD patients

Target Population: ESRD patients undergoing Continuous Ambulatory Peritoneal Dialysis (CAPD)

Inclusion Criteria

- ESRD patients who underwent CAPD catheter insertion at Dr. Saiful Anwar Hospital
- Treatment period: August 2014 – December 2024
- Declared deceased with CAPD-related peritonitis as the cause of mortality

Exclusion Criteria

- ESRD patients on CAPD who deceased due to causes other than peritonitis

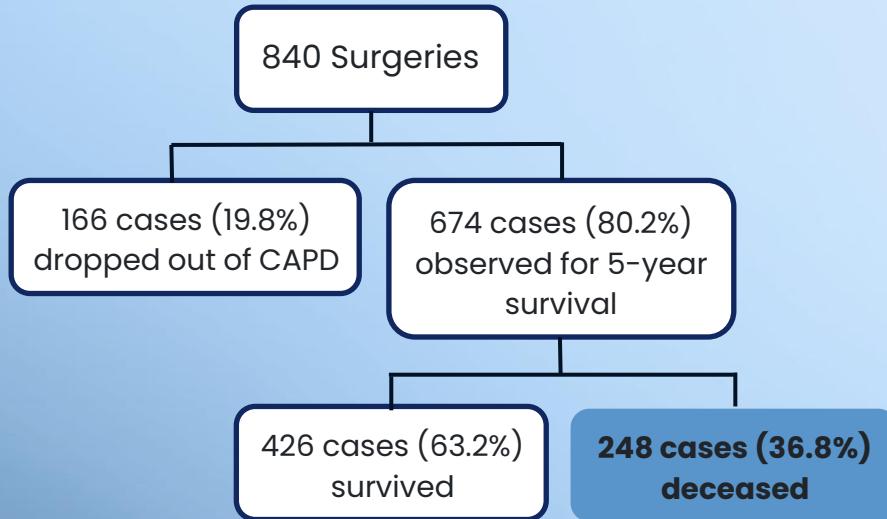
Diagnosis Criteria for CAPD-Related Peritonitis

Diagnosis is established if **at least two** of the following criteria are met:

1. Abdominal pain or cloudy peritoneal dialysate fluid
2. Leukocytosis in dialysate fluid (White blood cell count > 100 cells/ μL with $> 50\%$ PMN cells)
3. Identification of pathogenic bacteria by culture of peritoneal dialysate fluid

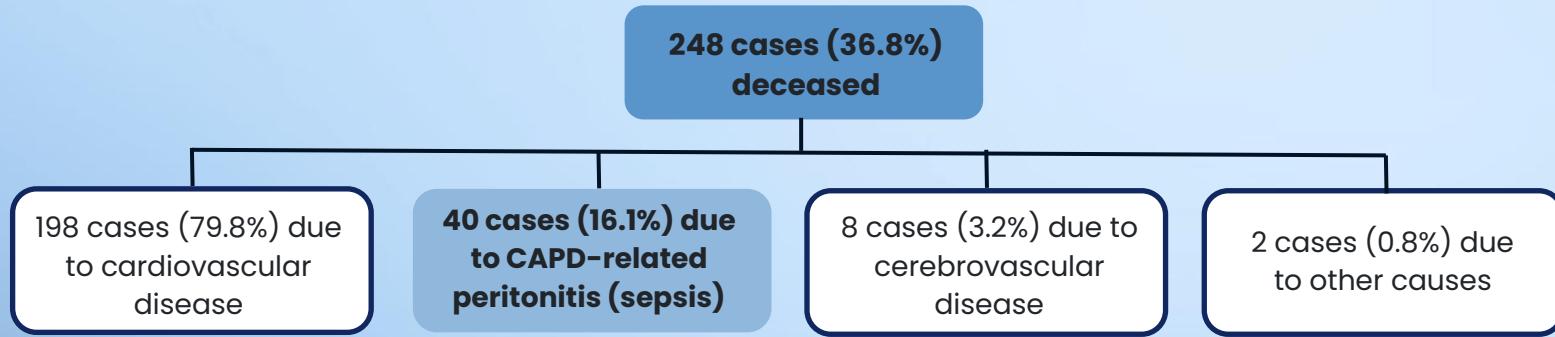
*Li PKT, Chow KM, Cho Y, et al. ISPD peritonitis guideline recommendations: 2022 update on prevention and treatment. *Peritoneal Dialysis International*. 2022;42(2):110-153

Results



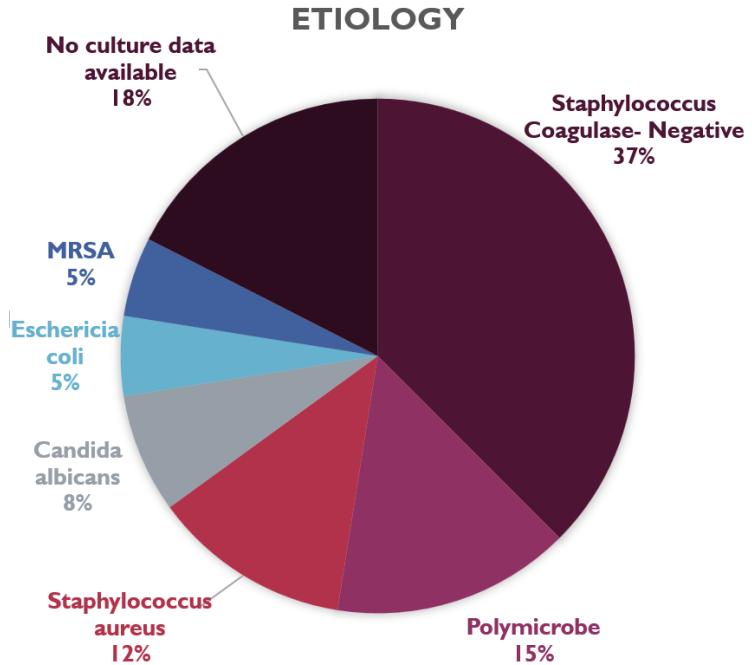
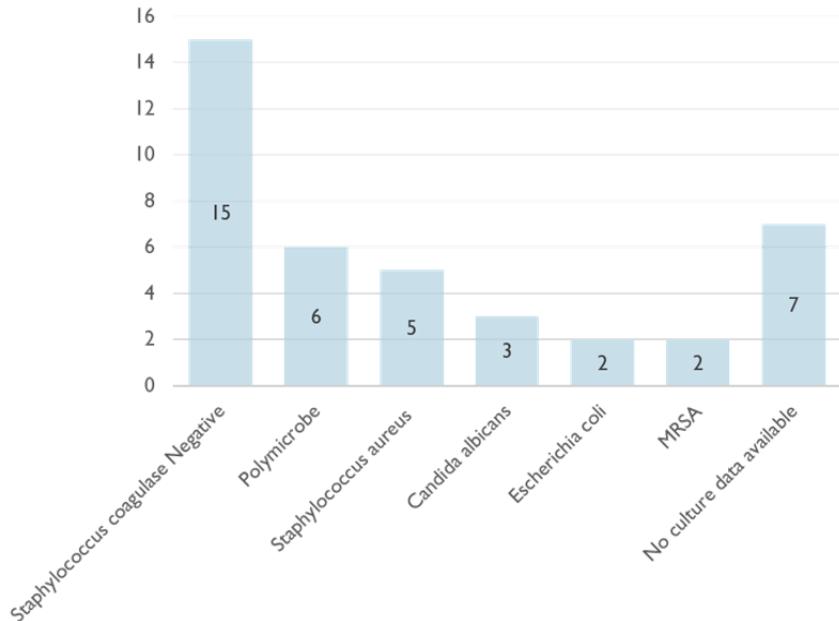
Characteristics	N=674	(%)
Gender		
Male	389	58.8
Female	285	41.2
Age, years		
<12	11	1.6
12-25	61	9.1
26-45	195	28.9
46-59	287	42.6
>60	120	17.8
Comorbid		
Hypertension	396	57.7
Hypertension and DMT2	278	42.3

Results



Results

Etiology of CAPD-related Peritonitis



Discussion

Staphylococcus Coagulase-Negative

- **Most common cause** of CAPD-related peritonitis in deceased patients → **37% cases**
 - **Similar findings** in Bach et al. (Vietnam), Nardelli et al. (Italy), Song et al. & Guo et al. (China) – Gram-positive, coagulase-negative *Staphylococcus* **up to 87%**
- Normal skin flora, mainly *S. epidermidis*, the most frequent PD-related pathogen.
 - Often causes **persistent and recurrent peritonitis** → **may require catheter removal and replacement**

Polymicrobial Infection

- Defined as **isolation of >1 organism** in a single peritonitis episode; more frequent in patients with prior peritonitis history
 - **Gram-positive** polymicrobial infections (~20%) → **better prognosis**, higher cure and catheter retention rates
 - **Gram-negative, anaerobic, or fungal infections** → **poorer outcomes**, higher rates of catheter removal and transfer to hemodialysis
- **15% of polymicrobial pathogens found in this study**
 - **Similar findings** in Bach et al. (Vietnam), 7 cases (8%) of CAPD peritonitis due to polymicrobial infection

1. Bach N, et al. *BMC Nephrol*. 2025;26(1).
2. Nardelli L, et al. *BMC Nephrol*. 2024;25(1):1-9.
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4. Guo S, et al. *Front Med (Lausanne)*. 2023;10:1132695.

Discussion

Staphylococcus aureus

- **5 cases (12 %) identified in this study**
 - Bach et al. (Vietnam): *S. aureus* was the most common coagulase-positive *Staphylococcus* causing CAPD peritonitis → **clinically severe peritonitis due to multiple virulence factors**
- **Associated with poor outcomes**
 - 20% recurrence rate
 - 23% catheter removal
 - 18% transfer to hemodialysis
 - 2% mortality

Candida albicans

- **3 cases (8%) identified in this study**
 - Fungal peritonitis incidence in CAPD ranges from <5%–15%; ***Candida* spp. cause >90% of episodes, mostly *C. albicans*.**
- **Difficult to treat effectively, often leading to:**
 - High catheter removal rates
 - Permanent transfer to hemodialysis
 - Poor patient outcomes

*ISPD guidelines recommend **immediate catheter removal** once fungal peritonitis is diagnosed, followed by **2 weeks of antifungal therapy***

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Discussion

Escherichia coli

- **2 cases (5%) identified in this study**
 - Bach et al. (Vietnam): E. coli caused 14 cases (8%) of CAPD peritonitis
 - Guo et al. (China): E. coli was the **most common multidrug-resistant organism (MDRO)**, responsible for 79 cases (54%)
- Clinical presentation **tends to be more severe**, with fever, abdominal pain, nausea, vomiting, and diarrhea
- **Prior antibiotic therapy** is a major risk factor for Gram-negative enteric peritonitis

Methicillin-Resistant Staphylococcus aureus (MRSA)

- **2 cases (5%) identified in this study.**
 - Guo et al. (China): MRSA caused 34 cases (23%) of CAPD peritonitis
- **MRSA is associated with worse outcomes compared to other multidrug-resistant S. aureus strains:**
 - Causes more severe peritonitis
 - Leads to longer hospital stays
 - Results in higher mortality rates

Rising antimicrobial resistance complicates empirical antibiotic selection, which should be adapted to local CAPD center conditions as recommended by ISPD guidelines

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Conclusions

The substantial mortality rate linked to CAPD peritonitis highlights the **critical need for enhanced infection prevention strategies and prompt targeted interventions**

Improving CAPD patient education, reinforcing aseptic techniques, and refining antibiotic stewardship programs are imperative to reducing peritonitis-related deaths and improving long-term patient outcomes.

Future research should focus on **developing predictive models for high-risk patients and evaluating the effectiveness of emerging antimicrobial therapies.**