

The Brazilian Journal of INFECTIOUS DISEASES

www.elsevier.com/locate/bjid



Clinical Image

Emphysematous cystitis: A rare form of urinary tract infection



Louise Cavalcanti Salles ⁽¹⁾ ^{a,*}, Tacilla Hanny de Souza Andrade ⁽¹⁾ ^a, Rafael Siqueira Athayde Lima ⁽¹⁾ ^b, Geraldo Bezerra da Silva Junior ⁽¹⁾ ^c

^a Hospital Geral de Fortaleza, Fortaleza, CE, Brazil

^b Hospital Universitário Walter Cantídio (EBSERH), Fortaleza, CE, Brazil

^c Universidade de Fortaleza, Centro de Ciências da Saúde, Curso de Medicina, Fortaleza, CE, Brazil

A R T I C L E I N F O

Article history: Received 21 September 2023 Accepted 31 October 2023 Available online 17 November 2023

A 74-year-old woman was admitted to the emergency department reporting dysuria and hematuria for 20 days. She had diabetes mellitus and recurrent Urinary Tract Infections (UTI), as well as cervical cancer treated with radiotherapy 25 years ago. Physical examination revealed hypotension (92×60 mmHg), heart rate 112 bpm, axillary temperature 36.7 °C, abdominal pain, drowsiness and disorientation. Laboratory findings evidenced anemia and leukocytosis (Hb 6.8 g/d, white blood count 16.800 mm³), c-reative protein 288 mg/dL, creatinine 3.0 mg/dL, urea 136 mg/dL. The urinary analysis showed leukocyturia, hematuria and bacteriuria. Due to urinary sepsis, antibiotic therapy (cephtriaxone) was started, and hemodialysis was required. She was admitted to the intensive care unit. Urine culture isolated Klebsiella pneumoniae (100.000 UFC/mL) ciprofloxacin and amoxicillin clavulanate resistant and sensitive to the other antibiotics tested. Abdominal tomography revealed gas in the interior and walls of the bladder. A new tomography was performed after 14-days, and significant radiological improvement was noted. Since she had been through prolonged hospitalization and multiple sepsis, she developed critical neuropathy and was transferred to a special care unit, dying after forty (40) days of hospitalization. She remained on dialysis therapy throughout the period (Figure 1).

Emphysematous Cystitis (EC) is a rare form of Urinary Tract Infection (UTI) characterized by gas formation on the bladder walls. The incidence rate of EC reported among diabetic women is 91.5 per 1000 person-years for UTIs in general.¹ The presentation includes: asymptomatic forms, classic cystitis up to sepsis.² The most common agents are *Escherichia* coli and *Klebsiella pneumoniae*.³ The main risk factors are Diabetes Mellitus (DM), female gender and neurogenic bladder.⁴ Clinical findings and imaging tests are important for the establishment of the correct diagnosis. Antibiotic therapy is the standard treatment. Prognosis is variable and early diagnosis favors better clinical evolution. Mortality varies between 7 %–10 %.^{2,5}

* Corresponding author.

E-mail address: louisecsalles@gmail.com (L.C. Salles).

https://doi.org/10.1016/j.bjid.2023.103700

^{1413-8670/© 2023} Sociedade Brasileira de Infectologia. Published by Elsevier España, S.L.U. This is an open access article under the CC BY-NC-ND license (http://creativecommons.org/licenses/by-nc-nd/4.0/)



Figure 1–(A) Abdominal tomography showing gas inside the bladder and in its walls, (B) New abdominal tomography 14-days later evidencing significant improvement.

Informed consent

Informed consent was obtained for publication of this case.

Authors' contribution

All authors contributed to data collection and discussed the final version of paper.

Conflicts of interest

The authors declare no conflicts of interest.

REFERENCES

- 1. Schicho A, Stroszczynski C, Wiggermann P. Emphysematous cystitis: mortality, risk factors, and pathogens of a rare disease. Clin Pract. 2017;7:930.
- 2. Thomas AA, Lane BR, Thomas AZ, Remer EM, Campbell SC, Shoskes DA. Emphysematous cystitis: a review of 135 cases. BJU Int. 2007;100:17–20.
- 3. Abusnina W, Shehata M, Nassri S, Zeid F. Emphysematous cystitis. Cleve Clin J Med. 2019;86:10–1.
- 4. Wang JH. Emphysematous cystitis. Urol Sci. 2010;21:185–6.
- Grupper M, Kravtsov A, Potasman I. Emphysematous cystitis: illustrative case report and review of the literature. Medicine. 2007;86:47–53.