

A pulmonary abscess due to *Klebsiella pneumoniae* carbapenemase 3

Maria Ana Canelas, Tatiana Fonseca

CASE REPORT

An 81-year-old male, with recurrent respiratory tract infections, and multiple antibiotherapy cycles in the previous three months, was admitted in the emergency department, referring fever and dyspnea for one week. Clinical, laboratory and imaging findings led to admission for pneumonia with respiratory failure. On the X-ray (Figure 1): “round hypotransparency, cavitated, in the right lower base”. For better characterization it was performed a chest CT scan (Figure 2) that revealed “bulky image, cavitated, with thick walls and polilobulated and heterogeneous, in the right inferior lobe, measuring 66x55 mm”. It was first initiated, empirically, clindamycin, for 20 days. Nevertheless, there was not improvement, and so, it was executed a bronchoscopy, allowing the isolation of *Klebsiella pneumoniae* carbapenemase (KPC) (clone ST 147) on the bronchoalveolar lavage. The antibiogram showed: sensitivity to amikacin; CIM Meropenem: ≤ 8 mg/L; antibiogram: < 2 mg/L; CIM colistin: ≤ 1.5 mg/L. He was then started on meropenem 2 g three-times per day and amikacin 500 mg twice per day, for 30 days, with improvement on the chest CT control (Figure 3), being discharged, after 97 days.

DISCUSSION

After an outbreak of KPC-3 (expressing the gene VIM-1) in our hospital, a protocol of treatment trial was

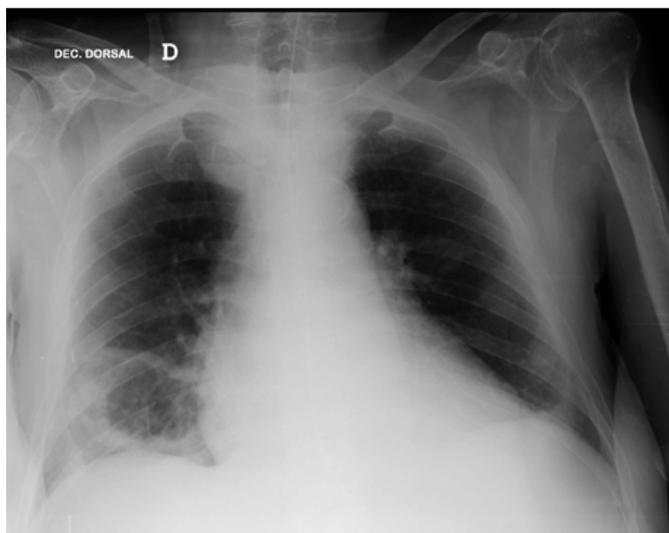


Figure 1: Round hypotransparency, cavitated, in the right lower base.

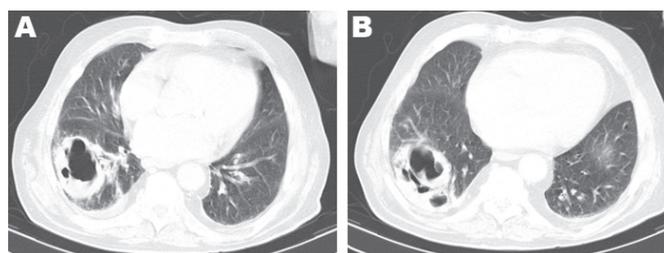


Figure 2: (A, B) “Bulky image, cavitated, with thick walls and polilobulated and heterogeneous, in the right inferior lobe, measuring 66x55 mm.”

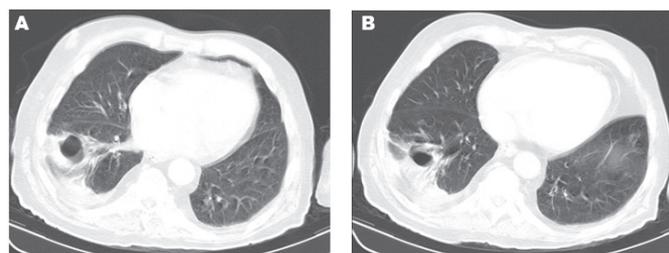


Figure 3: (A, B) Improvement on the chest computed tomography control.

Maria Ana Canelas¹, Tatiana Fonseca¹

Affiliations: ¹Internal Medicine Residents, Centro Hospitalar Vila Nova de Gaia/Espinho, Porto, Portugal.

Corresponding Author: Maria Ana Canelas, Internal Medicine Resident, Centro Hospitalar Vila Nova de Gaia/Espinho, Porto, Portugal; E-mail: anavc995@hotmail.com

Received: 22 June 2016

Accepted: 22 July 2016

Published: 16 September 2016

implemented, being of the most value the antibiogram. Misidentification of KPC is common with standard susceptibility testing. The most easily performed confirmatory test is the modified Hodge test, which has been found to be 100% sensitive, although not specific. Definite confirmation of KPC production requires molecular methods such as PCR [1]. There is some debate concerning the appropriate dosage and the most favorable pharmacokinetic/pharmacodynamic profiles in this cases [2, 3], but unfortunately the optimal treatment is unknown. The use of aminoglycosides, polymyxin combinations and tigecycline appeared to have higher success rates. Carbapenem and polymyxin monotherapy had much lower associated success rates [4]. Literature describes a wide range of approaches, from simple conservative treatments associated with specific antibiotic therapy, to surgical intervention, using endoscopic or percutaneous drainage [5]. Elores, a combination of ceftriaxone, disodium edetate and sulbactam, showed high susceptibility to KPC and Extended spectrum beta-lactamase (ESBL) producing pathogens, including the ones expressing VIM-1 [6]. The reason comes for the synergic activity from the Elores combination, with reports of efficacy and safety, and its use can be considered a drug of choice for treating KPC [7]. In our case, we report a more conservative treatment, with the use of high-dose of a carbapenem associated with an aminoglycoside, for a long period, with a favorable outcome. Furthermore, in our case, Elores could not be used, since disodium edetate it is not available in our country, and the antibiogram showed resistance to ceftriaxone.

CONCLUSION

The lack of published material and the emergency of these multiresistente microorganisms it is a challenge in the development of new treatment, control and prevention strategies. In the more serious cases of infections due to KPC, like abscesses, their adequate drainage, when possible, and the most appropriate antibiotic scheme showed a better outcome.

Keywords: Abscess, *Klebsiella pneumoniae* carbapenemase (KPC), Multiresistente microorganisms, Outbreak

How to cite this article

Canelas MA, Fonseca T. A pulmonary abscess due to *Klebsiella pneumoniae* carbapenemase 3. J Case Rep Images Med 2016;2:86–88.

Article ID: 100027Z09MC2016

doi:10.5348/Z09-2016-27-CL-20

Acknowledgements

All the staff members who were involved in the care of the patient.

Author Contributions

Maria Ana Canelas – Substantial contributions to conception and design, Acquisition of data, Analysis and interpretation of data, Drafting the article, Revising it critically for important intellectual content, Final approval of the version to be published

Tatiana Fonseca – Substantial contributions to conception and design, Acquisition of data, Analysis and interpretation of data, Drafting the article, Revising it critically for important intellectual content, Final approval of the version to be published

Guarantor

The corresponding author is the guarantor of submission.

Conflict of Interest

Authors declare no conflict of interest.

Copyright

© 2016 Maria Ana Canelas et al. This article is distributed under the terms of Creative Commons Attribution License which permits unrestricted use, distribution and reproduction in any medium provided the original author(s) and original publisher are properly credited. Please see the copyright policy on the journal website for more information.

REFERENCES

1. Arnold RS, Thom KA, Sharma S, Phillips M, Johnson JK, Morgan DJ. Emergence of *Klebsiella pneumoniae* Carbapenemase (KPC)-Producing Bacteria. Southern medical journal 2011;104(1):40–5.
2. Curcio D, Verde PE. Comment on: Efficacy and safety of tigecycline: A systematic review and meta-analysis. J Antimicrob Chemother 2011 Dec;66(12):2893–5.
3. Cunha BA. Pharmacokinetic considerations regarding tigecycline for multidrug-resistant (MDR) *Klebsiella pneumoniae* or MDR *Acinetobacter baumannii* urosepsis. J Clin Microbiol 2009 May;47(5):1613.
4. Hirsch EB, Tam VH. Detection and treatment options for *Klebsiella pneumoniae* carbapenemases (KPCs): An emerging cause of multidrug-resistant infection. J Antimicrob Chemother 2010 Jun;65(6):1119–25.
5. Di Carlo P, Pantuso G, Cusimano A, et al. Two cases of monomicrobial intraabdominal abscesses due to KPC–3 *Klebsiella pneumoniae* ST258 clone. BMC Gastroenterol 2011 Sep 30;11:103.
6. Chaudhary M, Payasi A. Molecular characterization and antimicrobial susceptibility study of *Acinetobacter*

- baumannii clinical isolates from Middle East, African and Indian patients. *Journal of Proteomics and Bioinformatics* 2012;5(11):265–9.
7. Gupta R. Antibiotic Adjuvant Therapy for Multi-Drug Resistant Carbapenemases Producing *Klebsiella pneumoniae* Associated Sepsis: A Case Study. *J Clin Diagn Res* 2016 Apr;10(4):DD08–9.

Access full text article on
other devices



Access PDF of article on
other devices

