Biomedical Laboratory Sciences

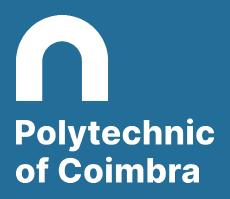
Beatriz Louçano

Clinical Case

Clinical Microbiology







Introduction

- Female;
- 32 years;
- Previously diagnosed with chronic kidney disease; •
- The patient was on dialysis through central venous catheter (CVC);



Two blood cultures collected from a peripheral vein to BACT/ALERT FA Plus blood culture bottles.

2º

Placed in the BactAlert Virtuo equipment (Biomérieux) , incubated for 5 days at 37°C. Monitors colorimetric changes indicated by the presence of CO2 = positive blood cultures.

Kim SC, et al. (2019); Kaase M, et al. (2009); Miller LM, et al. (2016);

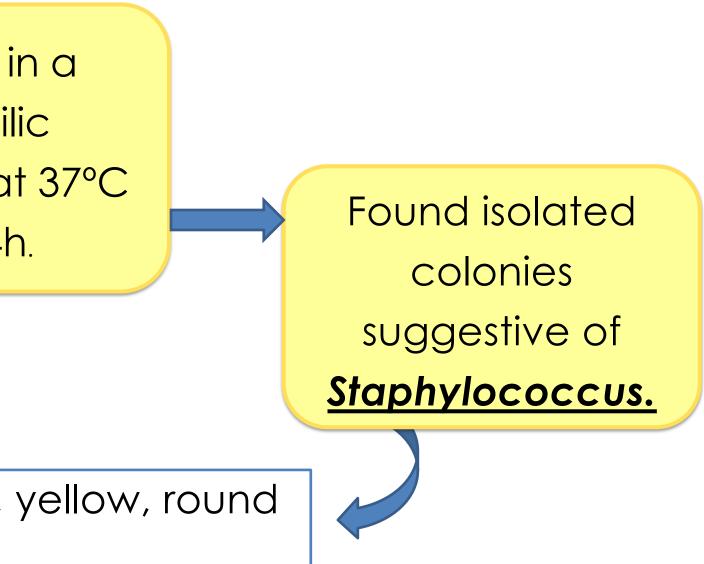
Nutrient medium with polymeric spheres adsorbing antibiotics, and a colorimetric sensor indicating the production of CO2.

The two blood cultures were positive and were transferred to Columbia Agar medium with 5% sheep blood.

Incubated in a capnophilic atmosphere at 37°C for 18/24h.

Gram positive bacterium with appearance of 33 small, yellow, round colonies

Kim SC, et al. (2019); Kaase M, et al. (2009); Miller LM, et al. (2016);



- The automated equipment, Bruker MALDI-Biotyper (Beckman Coulter), confirmed the presence of Staphylococcus aureus in the patient's sample.
- Antibiotic Susceptibility Testing (AST) were carried out semi-automatically using the **DxM MicroScan WalkAway** (Beckman Coulter) equipment.

Kim SC, et al. (2019); Kaase M, et al. (2009); Miller LM, et al. (2016);

Table 1 - Antibiotic Susceptibility Testing	
Isolated:	<u>Staphylococcus</u>
	<u>aureus</u>
Gentamicin	S
Trimethoprim	S
Vancomycin	S
Oxacillin	R
Penicillin G	R

Legend table 1 - Degrees of sensitivity:

S – Sensitive; **I** – Intermediate; **R** – Resistant

Table 2 - Screening for Carbapenemase-Producing Enterobacteriaceae

Results:	Gene
IMP1	Not detected
VIM	Not detected
NDM	Not detected
KPC	Detected
OXA48	Not detected

> Complementary results of biochemistry and hematology:

- Biochemical values are concordant with the previous diagnosis of CKD.
- **Protein C** and **procalcitonin** were increased \bullet
- In the blood count leukocytosis and neutrophilia, erythrocytes and hemoglobin ulletwere decreased, representative of anemia.

Discussion

\succ <u>Results from table 1:</u>

- There is resistance to oxalicin and penicillin G, and sensitivity to vancomycin, trimethoprim and gentamicin.
- The majority of Staphylococcus aureus are producers of penicillinases, which are specific β -lactamases to penicillin, causing destruction of the β -lactam ring.

Discussion

<u>Results from table 1:</u>

- The bacterium in question is resistant to oxacillin and penicillin G, which are from ulletthe family of penicillins and therefore we are in the presence of methicillin-resistant Staphylococcus aureus (MRSA).
- The most indicated antibiotic will be vancomycin. \bullet

Back KT, et al. (2014); Bonomo RA, et al. (2017); Hassoun A, et al. (2017); Hirao Y, et al. (2012); Miller LM, et al. (2016); Palavecino EL, (2020); Tjandra KC, et al. (2022);

Discussion

\succ <u>Results from table 2:</u>

- The patient had a positive KPC gene in the test for Carbapenemase-Producing Enterobacteriaceae (table 2), meaning that is colonized with KPC in her commensal flora.
- Should not be given β -lactamases antibiotics, because they destroy commensal flora allowing the growth of this opportunistic bacteria.
- The patient must comply with strict hygiene rules to prevent urinary infections or contaminating other patients with KPC.

the

Conclusion

Suspected diagnosis of : Sepsis crisis \bullet

The patient's final diagnosis: Persistent bacteremia (septicemia) caused by the ulletpresence of methicillin-resistant Staphylococcus aureus (MRSA) originated in the CVC. KPC gene is present in the patient's intestinal flora.

Bæk KT, et al.(2014); Bonomo RA, et al.(2017); Hassoun A, et al.(2017); Hirao Y, et al ,(2012); Miller LM, et al ,(2016); Palavecino EL,(2020); Tjandra KC, et al. (2022); Arnold RS, et al.(2011); Bassetti M, et al.(2020).

Bibliography

1. Arnold RS, Thom KA, Sharma S, Phillips M, Kristie Johnson J, Morgan DJ. Emergence of Klebsiella pneumoniae Carbapenemase (KPC)-Producing Bacteria. Southern medical journal. 2011;104(1):40-.

2. Bæk KT, Gründling A, Mogensen RG, Thøgersen L, Petersen A, Paulander W, et al. β-Lactam Resistance in Methicillin-Resistant Staphylococcus aureus USA300 Is Increased by Inactivation of the CIpXP Protease. Antimicrobial Agents and Chemotherapy. 2014;58(8):4593-.

3. Bassetti M, Peghin M. How to manage KPC infections. Therapeutic Advances in Infectious Disease. 2020;7.

4. Bonomo RA. β-Lactamases: A Focus on Current Challenges. Cold Spring Harbor Perspectives in Medicine. 2017;7(1).

5. Hassoun A, Linden PK, Friedman B. Incidence, prevalence, and management of MRSA bacteremia across patient populations-a review of recent developments in MRSA management and treatment. Critical care (London, England). 2017;21(1):211-.

6. Hirao Y, Ikeda-Dantsuji Y, Matsui H, Yoshida M, Hori S, Sunakawa K, et al. Low level β-lactamase production in methicillin-resistant staphylococcus aureus strains with β-lactam antibiotics-induced vancomycin resistance. BMC Microbiology. 2012;12(1):1-10.

7. Kaase M, Baars B, Friedrich S, Szabados F, Gatermann SG. Performance of MicroScan WalkAway and Vitek 2 for Detection of Oxacillin Resistance in a Set of Methicillin-Resistant Staphylococcus aureus Isolates with Diverse Genetic Backgrounds. Journal of Clinical Microbiology. 2009;47(8):2623-.

8. Kim SC, Lee S, Kim S, Cho OH, Park H, Yu SM. Comparison of Clinical Performance Between BacT/Alert Virtuo and BacT/Alert 3D Blood Culture Systems. Annals of Laboratory Medicine. 2019;39(3):278-.

9. Miller LM, Clark E, Dipchand C, Hiremath S, Kappel J, Kiaii M, et al. Hemodialysis Tunneled Catheter-Related Infections. Canadian Journal of Kidney Health and Disease. 2016;3(1).

10. Palavecino EL. Rapid Methods for Detection of MRSA in Clinical Specimens. Methods in molecular biology (Clifton, NJ). 2020;2069:29-45.

11. Tjandra KC, Ram-Mohan N, Abe R, Hashemi MM, Lee JH, Chin SM, et al. Diagnosis of Bloodstream Infections: An Evolution of Technologies towards Accurate and Rapid Identification and Antibiotic Susceptibility Testing. Antibiotics 2022, Vol 11, Page 511. 2022;11(4):511-.

Biomedical Laboratory Sciences

Beatriz Louçano

beatrizloucano@estescoimbra.pt





