

A Case of Infectious Endocarditis Due to *Gemella haemolysans* Complicated with Encephalomyelitis

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Abstract: *Gemella haemolysans* is one of cocci Gram-positive belonging to the family of Streptococcae. It is considered as commensal of the buccal mucous membrane, respiratory, digestive and genitourinary tractus. It is rarely involved in human pathology. We report a case of infectious endocarditis due to *Gemella haemolysans* complicated with encephalomyelitis in a 56-year-old patient without particular risk factor. The identification was done in the phoenix automaton BD and confirmed by the biochemical galleries. The evolution under Teicoplanin and after valvular replacement was favorable.

Keywords: *Gemella haemolysans*; infectious endocarditis; cocci.

INTRODUCTION

Gemella haemolysans is one of cocci Gram-positive aero-anaerobic optional belonging to the family of Streptococcae [1, 2], it is considered as a commensal species of the buccal mucous membrane, respiratory, digestive and genitourinary tractus [3, 4]. It puts difficulties of identification in the laboratory because of his fast discoloration in the Gram and of his similarity with the genre Streptococci. It may be complicated by encephalomyelitis [5, 6], of bone infection [7], meningoencephalitis [8]. However, the cases of endocardites were most reported [9, 10]. These infections concern as well the child as the adult. We report the case of an infectious endocarditis complicated with encephalomyelitis.

CASE REPORT

it concern H.F patient of 56 years old without particular pathological history who consulted for tumefaction of the straight forearm involving for two months in a context of anetous. An ultrasound doppler method allowed to highlight a false aneurysm of the ulnar artery so an surgical indication was decided. The cheking analysis before the surgy allowed to diagnose an organic renal insufficiency for which the patient benefited from a nephrostomy.

A bacteriological sample realized in the fall of the surgical procedure returned negative. In postoperative recoveries, the patient had installed in a rough way a left central Bell's palsy of type accompanied with a dysarthria. In the clinical examination, the patient had a kept, aware general

condition with a score of Glasgow in 15/15, a temperature in 38,5°C, a heart rate in 70 beatings by minutes. The muscle strength at the level of the lower and upper limbs was preserved, tendon reflexes were normal. The cardiac auscultation and above aortic trunks was unremarkable. The patient had no dental anomaly and did not undergo manipulation at this level recently. There was no notion of antibiotic therapy before the consultation. On the biological plan, the rate of C reactive protein (CRP) had passed of 76 mg / l to 142mg / l in 48 hours; The rate of leukocytes was 10 500/dl to 12000 / dl, the hemoglobin at 8,8g / dl and the platelets at 232 000 / dl.

A cerebral Scanner then a RMI realized found hypodense damage in the frontal zone in favour of an encephalomyelitis. The transthoracic ultrasound found vegetations on the ventricular hillside of the big bicuspid, mobile valve with a dilated left ventricle and a massive aortic insufficiency.

Giving these different finding, six bloodculture bottles were sowed, three in bottles BD Bactec anaerobic and three others in bottles BD Bactec aerobic. After 48 hours of incubation in the automaton Bactec 9240 of Becton Dickinson (BD), five bottles on six were indicated positive. The direct examination after centrifugation and Gram staining allowed to observe cocci grouped in the majority in heap with some in diplococcus, the Gram of which was weakly positive. Bottles were repiqued each on agar-agar columbia supplemented by 5 % of blood of sheep and on chocolate-brown agar-agar and incubated in the

sterilizing room under 5 % of CO₂. After 48 hours of incubation, it appeared small hemolytic colonies having the same aspect as the direct examination. These colonies were catalase denial, Resistance fighters in the optochine and do not degrade the esculine. The use of the automaton Phoenix 100 BD with galleries 77 had allowed the identification of *Gemella haemolysans* 99 %. the identification was confirmed by means of the gallery API ® 20 Strep system (BioMerieux).

The antibiogram had been realized at the same time by the method of distribution on chocolate agar according to the recommendations of the committee of the antibiogram of the French society of microbiology. Tree strains were inhibited by benzylpenicillin, aminopenicillins, the vancomycin, the teicoplanin, aminoglycosides, the acid fucidique, third generation cephalosporins and tetracycline. On the other hand, they were resistant in sulph drugs.

The initial treatment was benzylpenicillin which was secondarily changed by the teicoplanin seen that the patient presented a drug eruption.

The patient benefited from a replacement of the bicuspid aortic valve by a mechanical prosthesis and the closure of a perforing on the bicuspid valve. The evolution was favorable with one apyrexia obtained in 72 hours and a progressive reduction of CRP. The patient was declared outgoing after 6 weeks of hospitalization and kept a sequellaire Bell's palsy.

DISCUSSION

Initially classified in the genre *Neisseria* because of its fast discoloration in the gram, the knowledge of the structure of *Gemella haemolysans* allowed to classify it later among cocci Gram-positive [1, 11]. This genre also includes: *G. Morbillorum*, *G. bergeri*, *G. sanguinis*, *palaticanis G. and G. cuniculi* [4, 9]. Only *G. haemolysans* and *G. morbillorum* are recognized as being able to be pathogenic in humans. The infection caused by *Gemella haemolysans* arises generally on particular grounds: bad buccodental state, heart disease, immunosuppression, diverse prostheses [6, 7, 9, 12, 13]. However, some cases were reported at patient's without factors of particular risk [5, 14].

In this casereport, the patient had no underlying heart disease and no known risk factor. The clinical examination and in particular the buccodental examination did not allow to find a front door, the patient did not undergo dental manipulation. The endocarditis is the most frequent location (localization) of the infection to *Gemella haemolysans*, cases were reveled by some authors in Europe, in the United States and in Asia [9, 13]. Our case after reviewing the litterature is considered the 21th case of endocarditis to *Gemella haemolysans* documented in the world and the first case in the Maghreb. The endocarditis concerned or the bicuspid valve is the aortic valve [9], in this case

of species, it is about an infringement of the bicuspid valve.

The identification of *Gemella haemolysans* in the case of our patient was remitted in an automaton, then confirmed by the biochemical gallery API ® 20Strep. The sequencing of the ARNr 16S can be of a big contribution in the difficult cases [15]. *Gemella haemolysans* is sensitive in vitro in Benzylpenicillin, in cephalosporins, in glycopeptides and Resistance fighters in the trimethoprim-sulfamethoxazole [16].

Our patient evolved well under teicoplanin with an improvement on the clinical plan and the progressive reduction in the CRP. under the antibiotic therapy, all the cases of endocarditis reported *Gemella haemolysans* evolved well [13].

CONCLUSION

The infections to *Gemella haemolysans* are relatively rare but probably under diagnosed because of a slow growth and the similarity with the genre Streptococci. We report the first case of endocarditis to *haemolysans G.* in the Maghreb at a patient without any risk factor. The use of combined several tools of diagnosis in the laboratory will allow the isolation of this germ.

REFERENCES

1. Berger U. Aproposed new genus of gram-negative cocci: *gemella*. International bulletin of bacteriological nomenclature and taxonomy. 1961;11:17-19.
2. Stackebrandt E, Wittek B, Seewaldt E and Schleifer KH. Physiological, biochemical and phylogenetic studies on *gemella haemolysans*. Fems microbiology letters. 1982;13:361-365.
3. Heller D, Helmerhorst EJ, Gower AC, Siqueira WL, Paster BJ, Oppenheim FG. Molecular characterization of the microbial diversity in the early, *in vivo* formed, dental biofilm. Appl. environ. microbiol. Aem03984-1.
4. Berger U. Prevalence of *gemella haemolysans* on the pharyngeal mucosa of man. Medical microbiology and immunology. 1985;174:267-74.
5. Mneimneh S, Awada H, Shatila A. Achild with brain abscess due to *gemella haemolysans*. Ecpaediatrics. 2016;2(2):139-142.
6. Chinbo M, Addebbous A, Moutachakkir M, Rada N, Bouskraoui M, Chabaa L, Soraa N. Abcès cérébral à *gemella haemolysans* chez un enfant porteur d'une cardiopathie congénitale complexe. Ann biol clin. 2014;72(4):487-90.
7. Fangous MS. bone infections caused by *gemella haemolysans*. Med mal infect 2016.
8. Galen BT, Banach DB, Gitman MR, Trow TK. Meningoencephalitis due to *gemella haemolysans*. Journal of medical microbiology. 2014; 63:138-139.

9. Khan R, Urban C, Rubin D, Segal-maurer S. Subacute endocarditis caused by *Gemella haemolysans* and a review of the literature. *Scand J Infect Dis*. 2011;36:885-888.
10. Laudat P, Cosnay P, Icole B. Endocardite à *Gemella haemolysans* : une nouvelle observation. *Médecine et maladies infectieuses*. 1984;14:159-161.
11. Reyn A. Taxonomic position of *Neisseria haemolysans*. *International journal of systematic bacteriology*. 1970;20:19-22.
12. Barry R, Parminder JSJ, Anthony JS. *Gemella haemolysans* Infection in Total Hip Arthroplasty. *Case Reports in Orthopedics*, 2012.
13. Quaeset L. Endocardite à *Gemella haemolysans* chez un patient porteur d'une bioprothèse valvulaire aortique. *Med Mal Infect*, 2015.
14. Toshimasa H, Hideki U, Kunio Y, Yoshiyuki O, Hiroshi H, Norifumi T, Yoshihisa N. Recurrent *Gemella haemolysans* Meningitis in a Patient with Osteomyelitis of the Clivus. *Intern Med* 2013;52:2145-2147.
15. Woo PCY, Lau SKP, Fung AMY, Chiu SK, Yung RWH, Yuen KY. *Gemella* bacteraemia characterised by 16S ribosomal RNA gene sequencing. *J Clin Pathol*. 2003;56:690-693.
16. Buu-Hoi, Sapetra A, Branger C, Acar JF. Antimicrobial Susceptibility of *Gemella haemolysans* Isolated from Patients with Subacute Endocarditis. *Eur. J. Clin. Microbiol*. 1982;1:102-106.