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ARTICLE INFO

ABSTRACT

intravenous ceftriaxone.

Article history: Received 23 August 2020 Received in revised form 30 September 2020 Accepted 25 March 2021

Keywords: Leclercia adecarboxylata Endocarditis Opportunistic pathogen

Introduction

Leclercia adecarboxylata is a motile, gram negative bacillus in Enterobacteriaceae family [2]. The pathogen was first described by Leclerc as Escherichia adecarboxylata in 1962. In 1986 Leclercia, named in honor of Leclerc, was recognized as a new genus in the family Enterobacteriaceae [3]. The reclassification was enabled by the emergence of more sensitive testing methods such as DNA hybridization and computer identification studies [4]. The organism is widely distributed in nature, to include water and as a commensal in the gut [4]. Rarely, Leclercia is isolated from immunocompetent patients, but is primarily found to cause infection in the immunocompromised [5]. Due to limited reports of human infection, *L. adecarboxylata* was previously thought to be an opportunistic pathogen. However, this may reflect misdiagnosis, as the organism shares many biochemical features with Escherichia coli. As with other atypical species, mass spectrometry and automated identification systems have allowed more frequent and accurate identification [3]. Previous searches of the literature have only revealed two other reports of L. adecarboxylata endocarditis [6]. We report a case of L. adecarboxylata isolated from a Peripheral Inserted Central Catheter (PICC) and subsequent eustachian valve endocarditis.

Case report

A 62 year old Caucasian man with a history of three-vessel coronary artery bypass in 2016, sick sinus syndrome status post

* Corresponding author at: 1120 15th St, Augusta, 30912, Georgia. *E-mail address:* jfkidowu@gmail.com (F. Lawal). Micra pacemaker, hypertension, hyperlipidemia, colo-cutaneous fistula with colostomy bag and a ventral incision with wound us in vacuum-assisted closure (VAC), on Total Peripheral Nutrition (TPN) for bowel rest through a left arm PICC, presented to the Emergency

Leclercia adecarboxylata is a motile, gram negative bacillus in the Enterobacteriaceae family that is a rarely

isolated cause of disease, despite being ubiquitous in nature. A 2019 review article identified only 74

reported cases, most often in immunocompromised patients [1]. The organism is generally susceptible to

most antibiotics although multiantibiotic resistant strains have been reported. We report a case of a 62year-old Caucasian man with multiple co-morbidities treated for *L. adecarboxylata* endocarditis with

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vacuum-assisted closure (VAC), on Total Peripheral Nutrition (TPN) for bowel rest through a left arm PICC, presented to the Emergency Department with a 3 day history of progressively worsening dyspnea, high fever up to 39 °C, chills, vomiting, diaphoresis and generalized weakness. He had a previous history of three PICC associated blood stream infections, with his most recent infection in October 2019 due to *Staphylococcus epidermis*. He stated that his current episode felt like his previous bloodstream infections.

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On admission, he was febrile with a temperature 38.1 °C, tachycardic with heart rate of 130 beats per minute (bpm), hypotensive with blood pressure of 77/52 mmHg, and respiratory rate of 18 cycles per minute (cpm). Cardiac examination revealed a tachycardic rate of 130 s bpm with a regular rhythm and no murmur. Breath sounds were clear to auscultation and percussion bilaterally with no wheezes, rales or rhonchi. His abdomen was soft and nontender, and the wound VAC covering ventral incision present and ostomy bags covering enterocutaneous fistulae were clean and intact with no surrounding ervthema or induration. There were no signs of conjunctival hemorrhage, splinter hemorrhages, Janeway lesions, or Osler nodes. After administration of 2 L of normal saline, his heart rate improved to about 100 bpm, and the patient reported feeling significantly better. However, he still required norepinephrine 0.15mcg/kg/min to maintain a mean arterial pressure greater than 65 mmHg. The patient was started on intravenous (IV) vancomycin and cefepime as empiric antibiotic therapy for septic shock.

Laboratory findings showed white cell count of 6400/mm³, hemoglobin of 10.2 mg/dL, and platelet level of 76,000/mm³. Blood cultures were drawn from both forearm venipuncture and from the patient's PICC. Chest x-ray done at time of presentation was

http://dx.doi.org/10.1016/j.idcr.2021.e01091

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Table 1

Microbiology culture and susceptibility report.

	Leclercia adecarboxylata	
Drug	MINT	MDIL
Amoxicillin/Clavulanate	S	< = 2
Ampicillin	S	< = 2
Ampicillin/Sulbactam	S	< = 2
Cefazolin	S	< = 4
Cefepime	S	< = 1
Ceftriaxone	S	< = 1
Ciprofloxacin	S	< = 0.25
Ertapenem	S	< = 0.5
Gentamicin	S	< = 1
Imipenem	S	< = 0.25
Tobramycin	S	< = 1
Trimethoprim/Sulfamethoxazole	S	< = 20

normal, urine analysis was normal, and an abdominal/pelvis Computed tomography (CT) scan revealed no acute changes or signs of an infectious process. On day 2, two of four blood cultures drawn in aerobic and anaerobic bottles from the PICC was positive, showing gram negative rods on gram staining. The plate cultures grew two plus (++) gram-negative non lactose fermenting bacilli, that were later identified as *L. adecarboxylata* using, Matrix Assisted Laser Desorption/Ionization Time of Flight (MALDI-TOF), while blood cultures drawn from venipuncture sites were negative. The organism was pan-susceptible as shown in Table 1. The patient's PICC was removed to achieve source control, and IV vancomycin and cefepime were discontinued in favor of cefazolin 2 mg IV every 8 h based on sensitivities shown in Table 1.

A transthoracic echocardiogram (TTE) was performed to assess for valvular changes or vegetations to indicate endocarditis; however, the imaging study was limited by the patient's body habitus and was deemed poor quality. A subsequent transesophageal echocardiogram (TEE) revealed a "12 mm \times 4 mm mobile echodensity attached to the Eustachian valve" that was visualized moving freely and independently (Fig. 1). Upon review with the cardiologist, the valvular mass was visualized and confirmed to be freely mobile independent of the eustachian valve. The diagnosis of *L. adecarboxylata* infective endocarditis was subsequently made. A new PICC was placed after 6 days of negative blood cultures in preparation for outpatient parenteral antibiotic therapy (OPAT). His discharge antibiotic regimen was switched to IV ceftriaxone 2 g daily for 6 weeks for easy of dosing.



Fig. 1. Transesophageal echocardiogram. Arrow pointing to vegetation on Eustachian valve.

Discussion

Leclercia adecarboxylata is ubiquitous in the environment. It is found in food, water and other environmental sources [5]. However, it is rarely isolated from clinical samples and is not known as a frequent cause of infection. As such, it was initially thought to be an opportunistic pathogen in immunocompromised hosts [7]. This paucity of reported cases may reflect misdiagnosis at least in part, as the organism shares many biochemical features with *E. coli*. Advancements in microbiology techniques have led to accurate identification and segregation of *L. adecarboxylata* from *E. coli* and an increase in diagnosis [4]. These techniques include mass spectrometry and other automated identification systems [3]. In particular, Matrix Assisted Laser Desorption/Ionization Time of Flight (MALDI-TOF) mass spectrometry is preferred for precise species identification over conventional methods [3].

Since the first case reported in 1991, approximately 74 cases have been reported in English-language journals as of July 2020 [8]. Cases have been described both in the immunocompetent, and more frequently in the immunocompromised, with risk factors including central catheter use, malignancy, immunodeficiency, and invasive procedures [6]. In the immunocompetent, infections are primarily in the soft tissue and found in association with an aquatic environment [5]. It has also been isolated from a blood sample from a healthy donor [9], and implicated in several cases of bacteremia in the immunocompetent. In the immunocompromised, Leclercia adecarboxylata is implicated in cases which involve endocarditis [6], catheterrelated bacteremia [10], bacteremia and cellulitis [4] and spontaneous bacterial peritonitis [3]. Our patient was deemed to be relatively immunocompromised based upon his co-morbidities including gastrointestinal surgeries with dependence on TPN. Literature search revealed two other reports of L. adecarboxylata endocarditis [6,12]. The diagnosis of endocarditis was based upon the modified Duke criteria with one major criteria and three minor criteria (fever, predisposing heart condition, and positive blood cultures) [11]. As reports of *Leclercia adecarboxylata* endocarditis are rare, there are not established treatment regimens and antibiotic durations readily available [10]. As the organism isolated was pan-sensitive, we treated the patient with IV ceftriaxone 2 g daily for 6 weeks for ease of home dosing.

To conclude, we report a case of eustachian valve endocarditis caused by *Leclercia adecarboxylata*. Although it is quite rare, it may be an emerging cause of infection as advanced diagnostic technology allows its more frequent identification [3,4,7]. Clinicians should be aware of this pathogen, particularly in patients that are immunocompromised, have central catheters and pre-existing heart conditions.

Author contribution

Kashif Malik — case report design, writing. Ryann Davie — data collection, writing. Allison Withers — data collection, writing. Mohammad Faisal — data collection, writing. Folake Lawal — case report design, data collection, final review.

Funding statement

This research received no specific grant from any funding agency in the public, commercial or not-for-profit sectors.

Contributorship statement

All authors were involved in the design and conception of this manuscript.

Data sharing

All data pertaining to this research article are included within the manuscript as written.

Declaration of Competing Interest

The authors report no declarations of interest.

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