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Safety and efficacy profile of CSE-1034 as a prolonged de-escalation therapy in prosthetic joint infection: A case report

Abstract
Background

Although rare, infection is considered to be most dreadful of the prosthetic related complications resulting in repeated surgical intervention, extended hospitalization or sometimes in loss of implant or permanent disability if not treated promptly. Poor treatment outcome associated with prosthetic joint infections (PJIs) could be partly attributed to rise in anti-microbial resistance among the causative agents.

Case presentation

This is a first reported case of ceftriaxone + sulbactam + ethylenediaminetetraacetic acid (CSE 1034) being used as an de-escalation therapy for more than 24 days with good safety and efficacy outcome in a 78 year male patient with PJI associated with hip replacement surgery, treated initially with meropenem and colistin followed by prolonged de-escalation therapy (24 days).

Conclusions

In clinically complicated cases of deep infections where prolonged use of last resort antibiotics is used, CSE-1034 can be considered as a safe, efficacious and economical de-escalating antibiotic to complete the treatment course and prevent recurrence of infection, especially in PJI.

Consent Disclaimer:

As per international standard or university standard, patient's consent has been collected and preserved by the authors.

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32 **Introduction**

33 The number of hip replacement surgeries has rapidly increased in recent years. The joint
34 prosthesis is recognized as one of the most successful surgical interventions in medicine and
35 provides significant reduction in pain, improves joint function and minimizes disability^[1]. The
36 worldwide literature survey has shown infection rates for primary total hip revision to be 1-2%
37 and higher in cases undergoing revised total hip surgeries^{[2][3]}. Although rare, infections
38 following joint replacement surgery are the most devastating prosthetic-related complications
39 and challenging to treat ^[3].

40 Prevention of prosthetic joint infections is of utmost importance, and success in treating
41 these infections depends on extensive debridement, removal of dead and extraneous tissue and
42 most importantly effective antibiotic therapy ^[4]. Antibiotics form an important part of current
43 medical care and effective antibiotic prophylaxis constituting an important part in the treatment
44 of bacterial infections. Although infections following joint prosthetic surgeries are rare, however
45 with emerging bacterial resistance, its occurrence can be increasingly difficult to eradicate^{[5][6]}
46 PJIs with drug resistant pathogens may require prolonged medical management resulting in
47 extended and expensive hospital stays, repeated surgical intervention and sometimes can end up
48 in definitive loss of implant ^[5]. The failure of first choice of antibiotic used in empirical therapy
49 requires the treatment with second or third choice drugs that could be more expensive, less
50 effective and indeed more toxic. Here, researchers present a case of post-surgical infection
51 treated initially with a combination of meropenem and colistin followed by de-escalation therapy
52 of CSE-1034.

53 **Case presentation**

54 A 78-year-old male was admitted to intensive care unit department with chief complain
55 of oozing at operation site. He also complained of pain in left hip joint with instability while
56 walking.

57 The patient's medical history showed that he was operated for total left hip replacement
58 (THR) two months back. He also had a history of diabetes, hypertension and ischemic heart
59 disease. Moreover, the patient had also undergone coronary artery bypass grafting (CABG) about
60 8 years ago. After recently performed hip replacement surgery, the patient complained of
61 occasional pain. PBH X-ray had shown femur splinting in proximal medial aspect and the patient
62 was re-operated for tension wiring of implant and started with meropenem. On the post operation
63 third day patient had an episode of rigor; the patient was transferred from surgery ward to
64 intensive care unit department, for further treatment and management. Physical examination
65 revealed the patient to be afebrile with all his vitals normal, temperature: 98.4°F, blood pressure:
66 120/90 mmHg and pulse: 78 beats per min. Systemic examination of central nervous system
67 showed the patient was conscious and oriented, cardiovascular examination revealed S1 and S2
68 within normal limits, per abdomen was soft and non-tender and respiratory system was clear.
69 Hematology tests done revealed deranged TLC count (18500/cu-mm) with neutrophil count on
70 higher side (6.58×10^3 cells/cu-mm), deranged hemoglobin (11.8 g/dl); raised ESR (45 mm/hr.)
71 and hs-CRP (10.4 mg/dL) levels. Venous Doppler Ultrasonography of both lower limbs showed
72 normal flow in both lower limbs veins with no deep vein thrombosis. Mild sub-cutaneous edema
73 was observed in left leg on both sides and continuous discharge from the operation site was
74 noticed. Because of continuous discharge from wound; surgical site wash, wound exploration
75 and debridement of all deep infected tissues above fascia and below deep fascia were performed.
76 In the process, pulse lavage of the wound was done with 3-4 liters of normal saline. After wound
77 cleansing, closure was done in layers followed by dressing. Wound discharge was forwarded to
78 laboratory for culture and sensitivity testing. Based on Musculoskeletal Infection Society criteria,
79 a provisional diagnosis of PJI was arrived at, and the patient was re-shifted to ICU and started
80 immediately with intravenous (IV) dose of antibiotics (meropenem + colistin) and other
81 supportive treatment. The antibiotic choice was based on the hospital antibiogram data and the
82 patient's hematological parameters. Laboratory culture and sensitivity report didn't reveal any
83 pathogenic growth. After 48hr of ongoing antibiotic treatment, the patient was observed to
84 respond to treatment and the exudates from the surgical site started decreasing. The patient
85 continued to receive same IV treatment with antibiotics for 7 days. Repeat hematological tests
86 and other parameters confirmed patient's improvement. On 8th day of admission, the patient was
87 shifted to de-escalation therapy of CSE-1034 (3gm/12hr), considering its broad-spectrum activity

88 in targeting various resistant mechanisms of pathogens. The patient responded well to the de-
89 escalation treatment and was thus continued with CSE-1034. After 10 days of treatment, the
90 patient was hemodynamically stable and shifted to ward. He was discharged on 18th day of
91 admission and advised to continue CSE-1034 1.5 gm /12hr via IV and other basic supportive
92 medicine for 2 weeks. The patient was advised for follow-up and to report immediately in case
93 of fever, pain at operation site, convulsions, headache, chest pain and breathlessness and loss of
94 consciousness. On regular follow-ups and clinical examination, it was observed that the patient
95 didn't show any sign and symptoms of recurrence or superinfection and was hemodynamically
96 stable.

97 **Discussion**

98 Despite the high success rate, joint prosthetic surgeries are not without complications. Of
99 all prosthetic related complications, infection is probably the most threatening one. Although
100 rare, prosthetic infections after total hip replacement surgeries require prolonged surgical and
101 medical management. The costs of treating an infection after total hip replacement are reported
102 to a minimum amount of 50,000 US dollars per patient. Moreover, the steep rise in the number of
103 operations for implanting these prostheses in past few years, has been simultaneously
104 accompanied by number of PJIs ^[7]. Although, the recent technical advances in the field of
105 medicine accompanied by use of laminar air flow, exhaust systems, antibiotic-loaded acrylic
106 bone cement and antibiotics have all contributed to reduced infection rate, however PJIs still
107 occur in 1-3% of patients ^[7].

108 Frequently, the organisms implicated in infected prosthetic joints are usually relatively
109 benign organisms or simple contaminants. These microorganisms may penetrate wound during
110 surgical procedures from both endogenous and exogenous sources including patient's external
111 microbiota, microbiota of surrounding surgical team, hospital environment, surgical instruments
112 and even contaminated implants. PJIs that develop during first year after prosthetic implant are
113 considered to be SSIs and are usually treated using broad spectrum antibiotics. Though broad
114 spectrum, anti-microbial therapy at the time of induction is given, that helps to cut the risk of
115 infection at the surgical site ^[8]. These benign microorganisms tend to become pathogenic taking
116 advantage of suppressed immune system, associated co-morbidities along with sterile surgical
117 sites making certain individuals more vulnerable to infections.

118 Here, researchers present a case report of PJI treated successfully with initial therapy of
119 meropenem + colistin, and de-escalation therapy of CSE-1034. The patient responded well to the
120 treatment and was discharged on 18th day of admission with the advice to continue CSE-1034 for
121 2 weeks. Hence the present report highlights the importance of CSE-1034 in deep infections
122 usually treated with only last resort antibiotics. The normal course of antibiotic treatment for PJIs
123 extends from 4-6 weeks. Moreover, drug induced thrombocytopenia is reported in patients
124 undergoing meropenem treatment for more than 10 days^[9]. Thus, in deep infection cases where
125 carbapenems are used empirically and the treatment duration extends from 4-6 weeks, CSE-1034
126 can be used as deescalating antibiotic to complete the treatment course and cure the infection
127 without observing any side effects associated with prolonged meropenem therapy and
128 compromising the safety of patients. In support of our outcome, various studies in the past have
129 documented CSE-1034 as an effective treatment for MDR bacterial infections alone or as
130 combination therapy with colistin^{[10][11][12]}.

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132 **Conclusion**

133 This unique case study highlights the safety and efficacy profile of CSE-1034 in prolonged
134 duration treatment modules along with dramatic reduction in treatment-related costs as a de-
135 escalation therapy in treating PJI.

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