

A case of Snakebite complicated by *Morganella morganii* subspecies *morganii* Biogroup I infection

C Valsan, T Rao, A Sathiavathy

Citation

C Valsan, T Rao, A Sathiavathy. *A case of Snakebite complicated by Morganella morganii subspecies morganii Biogroup I infection*. The Internet Journal of Infectious Diseases. 2007 Volume 6 Number 2.

Abstract

A case of *Morganella morganii* subspecies *morganii* Biogroup I infection is confirmed by repeated isolation. Earlier patient failed to respond to empirical prescriptions of antibiotics, but promptly responded to a course of Cefotaxime to which the strain was sensitive. Snake bite wounds are secondarily infected with varied groups of organisms and specific isolation and identification is important to administer optimal antimicrobial therapy to avoid long term complications.

INTRODUCTION

Snake bite carries the consequences of envenomation primarily but also can produce a lesion at the bite site with extensive necrosis. The dead tissue can secondarily get infected by bacteria coming from the snake's mouth that may be inoculated at the moment of bite (1). Till recently very little was known about the bacteria responsible for the infection. In the recent past *Morganella morganii* was identified as an important pathogen associated with snake bite apart from the other hospital infections. *Morganella morganii* is a gram negative bacilli belonging to the family Enterobacteriaceae found in the environment and also in the intestinal tracts of man, mammals and the oral cavity of reptiles as a part of normal flora (1,2). Despite its widespread distribution it is an uncommon cause of community acquired infection and is most often encountered in postoperative and other nosocomial settings where it can cause urinary tract infections, sepsis, pneumonia, wound infection, chorioamnionitis, empyema and peritonitis. Here we present a case report of soft tissue infection with *M.morganii* following snake bite.

CASE REPORT

A 38 year old female was admitted to the surgical casualty ward after being bitten by a suspected poisonous snake on the right thumb. Patient was given anti-snake venom and prophylactic antibiotics (Ampicillin + Cloxacillin 500 mg IV 8th hourly) along with other first aid measures.

On the 3rd day the patient developed cellulitis with an abscess on the dorsum of the hand. Abscess was drained and

patient was put on a changed regime of broader spectrum antibiotics (Amoxicillin + Clavulanic acid) to cover the beta lactamase producing strains. But the abscess recurred and the patient had to undergo repeated incision and drainage. The direct smear made from these pus samples showed plenty of pus cells with gram negative bacilli on gram staining.

All the samples on culture grew *Morganella morganii* subspecies *morgani* biogroup 1 (3) with the same antimicrobial sensitivity-resistant to Ampicillin and 1st and 2nd generation Cephalosporins, but found to be sensitive to 3rd and 4th generation Cephalosporins, Aminoglycosides, Fluoroquinolones, Piperacillin and Tetracycline. After the second incision and drainage she was changed to Cefotaxime as guided by the antibiogram to which she responded and made a good recovery without any disability.

DISCUSSION

Soft tissue infections can be an important complication of snakebite with local envenoming. Various origins of bacteria at the site of venomous snakebite have been considered. Culture of fang, fang sheath and venom of snakes like Bothrops, Vipers, Rattlesnake and *Naja naja* had shown heavy colonization with many bacteria (2,4,5). These include Enterobacteriaceae like *Morganella* spp. and *E.coli*, Group D Streptococci, and anaerobes like Clostridia spp. In these cases the bacteria isolated were similar to that found in the abscesses from the bitten patients.

Other sources suggested include the victim's cloths and skin which were pierced by the fangs of the snake, or other

unsterile substances applied to the site of bite. In hospitalized patients the bacteria from the hospital could contaminate the lesion and cause nosocomial infection. In one prominent study *M. morganii* was isolated in 50% of abscesses occurring at the site of Bothrops bite. In another 12 year study *M. morganii* is reported as causing 10% of secondary infections following snake bite (4).

M. morganii is found to be inherently resistant to several antibiotics including Ampicillin, first and second generation Cephalosporins and to combinations of Amoxycillin and Clavulanic acid (6). It proved true in the present case as the patient did not respond to the initial antibiotics prescribed on empirical basis. The septic complications of snakebites need an optimal microbiological evaluation as varied groups of bacteria with unpredictable antibiotic sensitivity patterns has been reported.

The present case report enlightens the role of bacteriological culturing of septic materials from primarily and secondarily infected snake bite wounds and starting on appropriate

antibiotics at the earliest to prevent the disabling complications.

References

1. Jorge MT, Ribeiro LA . Infections in the bite site after envenoming by snakes of Bothrops genus. J.Venom.Anim.Toxins 1997; 3(2) Botucatu
2. Jorge T, de Mendonca JS, Ribeiro LA, da Silva ML, Kusano EJ., Codeiro CL. Bacterial flora of the oral cavity, fangs and venoms of Bothrops jararacca: possible source of infection at the site of bite . Rev Inst Med Trop Sao Paulo 1990; 32(1): 6-10.
3. Koneman EW, Allen SD, Janda WM, Schreckenberger FC. Enterobacteriaceae. Schreckenberger FC, Washington C Winn,Jr (eds).Colour atlas and textbook of diagnostic microbiology, fifth edition: page213, 222-223. Lippincott William and Wilkins 1997.
4. Avila -aguero ML, Valverde K, Gutierrez J, Parris MM, Faengezicht. Venomous snakebite in children and adolescents: A12 year retrospective study. J Venom Anim Toxin 7(1) Botucatu 2001
5. Ciszowski K, Hartwich A. Envenoming by Malayan Cobra (*Naja naja sputatrix*- case report). Przegl/ Lek 2004; 61(4): 421-6
6. Stock I, Wiedemann B. Identification and natural antibiotic susceptibility of *Morganella morganii*. Diagn Microbiol Infect Dis 1998; 30: 153- 65.

Author Information

Chithra Valsan, MD

Assistant Professor, Department of Microbiology, Jubilee Mission Medical College and Research Institute

T.V. Rao

Professor and Head of Department, Department of Microbiology, Jubilee Mission Medical College and Research Institute

A. Sathiavathy

Professor, Department of Microbiology, Jubilee Mission Medical College and Research Institute