A Case of Pneumonia Caused by *Ewingella americana* in a Patient with Chronic Renal Failure

Though the pathogenic significance and the reservoir of *Ewingella americana* have not been clarified, this organism has caused several pathogenic infections, especially in immunocompromised patients. We report a pneumonia in a patient with chronic renal failure, who had chronic rejection of transplanted kidney. *E. americana* was identified to be the pathogen of pneumonia with clinical symptoms and signs and radiological examination. As soon as he was treated with ceftriaxone and isepamicin, clinical improvement was followed with no further growth of *E. americana* or other pathogenic isolates from sputum culture. This suggests to be the case of pneumonia caused by *E. americana* for the first time in the Korean literature.

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**INTRODUCTION**

*Ewingella americana* is the only species of the genus of *Ewingella* in the family Enterobacteriaceae, first described from clinical specimens in 1983 (1). The pathogenic significance and niches of the reservoir have not been clarified. This organism rarely causes human infections and has been identified from various clinical specimens including wound, sputum, urine, stool, blood (1, 2), conjunctiva (3) and peritoneal dialysate (4). We present a chronic renal failure patient with fever and haziness in right lung field on chest radiography. Since the blood and urine cultures showed no other bacterial growth, pneumonia caused by *E. americana* suspected to be the origin of fever. Ceftriaxone and isepamicin (Yoohan, Seoul, Korea) were administered and the clinical and radiological findings were improved. As far as we know, this is the first report of a pneumonia caused by *E. americana*.

**CASE REPORT**

A 35-yr-old male who had undergone renal transplantation was admitted to the University Hospital of Keimyung for chronic rejection of the transplanted kidney on 1 May 2003. He had chronic renal failure for 7 yr. On the day of admission, peritoneal dialysis was initiated. After three days, he developed fever with right pleuritic chest pain. Physical examination revealed 38.2°C of fever and pale conjunctiva with pit-
itin, ceftazidime, ciprofloxacin, gentamicin, imipenem, piperacillin, piperacillin-tazobactam, and tetracycline, but only resistant to cephalothin (Table 2).

On 12 May 2003, subsequent sputum culture failed to grow either *E. americana* or any pathogenic bacteria. Chest radiography findings were improved as the clinical signs and symptoms disappeared. Ceftriaxone and isepamicin were administered for an additional week. And he was discharged with peroral medication of the antibiotics after the 15th day of admission.

**DISCUSSION**

Clinical infections due to *E. americana*, a rare Gram-negative bacilli with low pathogenic potential, have usually been involved in immunocompromised patients and caused peritonitis (4) and bacteremia (6, 7). Colonization in sputum (1), wound (8), and pseudobacteremia (9, 10) were reported in patients without clinical infections and in previously healthy patients. Sputum was reported to be the most common source of isolation (1, 2). Grimont reported that pneumonia or influenza has been developed only in 2 cases out of 5 strains (1), and

Farmer et al. listed 14 sputum isolates without clinical reviews (2). Therefore, the clinical significance of the isolate has not been clear due to no further case studies. Bacteremia was the most common infections of *E. americana* (6, 7, 11, 12). Very recently, Tsokos reported a patient who died of Waterhouse-Friderichen syndrome had growth of *E. americana* in heart and spleen blood samples of postmortem bacteriologic cultures and stated that the pathogenic potentiality of this strain (13).

Though several cases were published in the literature, little is known about the natural habitat of organism. However, McNeil proposed that the citrate solutions prepared in the hospital for coagulations study may have constituted an inanitated environmental reservoir and resulted in recurrent pseudobacteremia caused by *E. americana* (10). Also, the sources of infection considered to be the domestic water in peritonitis of a patient undergone peritoneal dialysis (4) and the contamination of ice bath as a probable cause of bacteremia (12).

On the other hand, Maertens et al. speculated on the contamination of catheter or inadequate hand hygiene as the source of infection (11).

*E. americana* can survive in water and grow preferentially at 4°C (10) and this isolate is not known to be the normal flora of respiratory tract. Since this study did not include evaluations dealing with water or other environmental sources throughout the admission, we could only presumed that the general hygiene or water contamination might have played a role in this infection.

According to the overall considerations of symptoms and signs, laboratory findings and the clinical course following antibiotics administration, we conclude that *E. americana* isolated in sputum could be the pathogen of pneumonia in this patient. Therefore we present the first case, to our knowledge, of *E. americana* pneumonia in a chronic renal failure patient. We have neither identified *E. americana* before in our laboratory nor isolated since then from the specimens of other patients. Regarding to the host and pathogenic potentiality, *E. ameri-

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**Table 1. Biochemical characteristics of *E. americana* strains**

| Reactions                  | % of positive reaction* | Kati† | The present strain |
|----------------------------|-------------------------|-------|--------------------|
| Indole production          | 0                       | not tested | -                 |
| Citrate (Simmons)          | 95                      | +     | +                 |
| Hydrogen sulfide (TSI)     | 0                       | -     | -                 |
| Urea hydrolysis            | 0                       | -     | -                 |
| Lysine decarboxylase       | 0                       | -     | -                 |
| Arginine dihydrolase       | 0                       | -     | -                 |
| Ornithine decarboxylase    | 0                       | -     | -                 |
| Motility (36°C)            | 60                      | -     | -                 |
| Malonate utilization       | 0                       | -     | -                 |
| D-Glucose, acid            | 100                     | +     | +                 |
| D-Glucose, gas             | 0                       | -     | -                 |
| Lactose fermentation       | 70                      | +     | +                 |
| Sucrose fermentation       | 0                       | -     | -                 |
| D-Mannitol fermentation    | 100                     | +     | +                 |
| Adonitol fermentation      | 0                       | -     | -                 |
| myo-Inositol fermentation  | 0                       | -     | -                 |
| D-Sorbitol fermentation    | 0                       | -     | -                 |
| L-Arabinoose fermentation  | 0                       | -     | -                 |
| Raffinose fermentation     | 0                       | -     | -                 |
| L-Phrarnose fermentation   | 23                      | -     | -                 |
| Maltose fermentation       | 16                      | +     | +                 |
| D-Xylose fermentation      | 13                      | -     | -                 |
| Esculin hydrolysis         | 50                      | +     | +                 |
| Acetate utilization        | 10                      | -     | -                 |
| Nitrate reduction          | 97                      | not tested | +              |
| Oxidase, Kovacs            | 0                       | -     | -                 |
| ONPG test                  | 85                      | -     | -                 |
| Yellow pigmentation        | 0                       | -     | -                 |

*Reactions of the published strains (2), †Results from Kati et al. (4).*

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**Table 2. Antimicrobial susceptibility results of *E. americana***

| Antimicrobial agents       | Interpretation |
|----------------------------|----------------|
| Amikacin                   | S              | S              |
| Ampicillin                 | S              | S              |
| Ampicillin-sulbactam       | NT†            | S              |
| Aztreonam                  | NT             | S              |
| Cefepime                   | S              | S              |
| Cefotaxime                 | S              | S              |
| Cefoxitin                  | NT             | S              |
| Cefalothin                 | R              | R              |
| Ciprofloxacin (ofloxacin-Katti) | S | S |
| Gentamicin                 | S              | S              |
| Imipenem                   | NT             | S              |
| Piperacillin               | NT             | S              |
| Piperacillin-tazobactam    | NT             | S              |
| Tetracycline               | NT             | S              |

*Results from Kati et al. (4), †NT, not tested.*
Pneumonia by *Ewingella americana* in Chronic Renal Failure

can become a rare but opportunistic pathogen of clinical infections, especially in an immunocompromised person. Even the etiology and route of infection have not been clearly stated, this organism, *E. americana*, has finally resulted in pneumonia and treated well with immediate usage of antibiotics in this patient. However, more cases and informations are needed to clarify and postulate the ecology and pathogenicity of *E. americana* in the hospital settings.

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