

## Case Report

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# Bacteremia caused by *Comamonas testosteroni* an unusual pathogen

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## Abstract:

*Comamonas* species are rare isolates in microbiology laboratories and have been infrequently reported as an infectious agent in routine clinical practice. They have a wide range of natural habitats including water, soil, and plants as well as from some hospital devices, such as intravenous lines and the reservoir water in the humidifiers of respiratory therapy equipment. *Comamonas testosteroni* is rarely recognized as a human pathogen. In spite of its uncommon human pathogenesis, there are few reports where it was reported as an aggressive opportunistic pathogen, and that was mostly related to *Testosterone* species. Herewith, we are reporting this pathogen from the blood of an immunocompetent female. The aim of this case report is to alert clinicians and laboratory physicians for the potential diagnosis and clinical approach of bloodstream infections caused by such unusual pathogens. This is the first documented case of bacteremia caused by *C. testosteroni* from India.

## Key words:

Bacteremia, *Comamonas testosteroni*, immunocompetent

## Introduction

*Comamonas testosteroni* is an aerobic Gram-negative bacillus that is oxidase positive, motile, nonspore forming, nonglucose fermenting, and arranged either singly or in pairs.<sup>[1]</sup> It was formerly known as *Pseudomonas testosteroni*, and now this bacillus was reclassified by molecular genetics. It was renamed as *C. testosteroni* as it is known to use testosterone as the sole source of carbon for their growth. Although it has low virulence potency, however, there are few case reports on its involvement in human infections. became clinically important after 1987 when reports began accumulating on human infections such as peritonitis, cellulitis, endocarditis, meningitis, pneumonia, endophthalmitis, and tenosynovitis<sup>[2]</sup> but cases of bloodstream infections caused by *C. testosteroni* have been infrequently reported. *C. testosteroni* is commonly found

in soil, water saprophytes, and plants. It has also been detected in the water of in-use hospital oxygen humidifier reservoirs, hemodialysis solutions, urine, mucus, prostate tissue, stool, and respiratory secretions of cystic fibrosis patients.<sup>[3]</sup> However, it has not been recognized as a component of the endogenous human microflora. The genus *Comamonas* originally contained following species: *Comamonas terrigena*, *C. testosteroni*, *Comamonas denitrificans*, *Comamonas nitrivorans*, and *Comamonas kerstersii*. It now contains 17 species, while *Comamonas acidovorans* has been separated from the genus on the basis of its 16S rRNA gene and is now known as *Delftia acidovorans*. Although pathogenic potential of *C. testosteroni* has not been studied well, it is usually considered as a commensals. Around 36 cases have been reported in medical literature concerning *C. testosteroni* infection so far.<sup>[3]</sup> Herewith, we are reporting the first documented case of *C. testosteroni* bloodstream infection from India.

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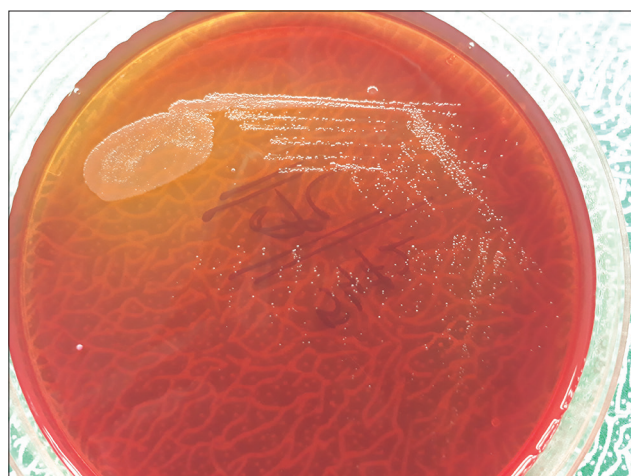
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## Case Report

A 46-year-old female from rural area admitted to our hospital with chief complains of high-grade fever (103°F), headache, fast breathing, and burning micturition for the past 6 days. On examination, other findings were such as respiratory rate 82/min, bilateral fine crepitation, pulse rate 146/min, and blood pressure 140/90 mmHg. Central nervous system, cardiovascular system, and per abdominal examinations were within normal limit. Other relevant laboratory findings include hemoglobin 10 gm%, total white blood cell count 24,200/cmm, neutrophil 90%, lymphocyte 10%, fasting blood sugar – 88 mg/dl, erythrocyte sedimentation rate 50 mm after 1 h, Na – 123 m. mol/L, K – 4.2 m. mol/L, blood urea – 45 mg/dl, serum creatinine – 1.8 mg/dl, urine routine microscopy shows pus cells 6–10/hpf, albumin-trace, and no parasites. Serological tests such as HBsAg, HIV, HCV, and dengue (NS1Ag, IgM, and IgG) were negative by Enzyme-linked immunosorbent assay. Chest X-ray shows bilateral opacities in lower lobe and ultrasound sonography (USG) abdomen was normal. Blood was collected with complete aseptic precaution in aerobic and anaerobic blood culture bottle (BacT/ALERT/3D; bioMerieux, Marcy l'Etoile, France). The patient was then started empirically with piperacillin + tazobactam and vancomycin. Aerobic blood culture bottle showed a positive sign of growth after 72 h. The broth was then subcultured on 5% sheep blood agar and MacConkey agar. After overnight incubation, MacConkey agar showed nonlactose fermenting colony but colony morphology was better seen after 48 h of aerobic incubation. Colonies were 1–2 mm in diameter, translucent, smooth surface, and nonpigmented [Figure 1]. Blood agar plate showed similar colony morphology and nonpigmented. Gram-stain was done from both the culture plates, and it was Gram-negative bacilli. It was motile by hanging



**Figure 1:** Nonlactose fermenting colony of *Comamonas testosteroni* on MacConkey agar

drop method. Routine biochemical tests were done and it was catalase positive, oxidase positive, indole not produced, urea not hydrolyzed, citrate not utilized, TSI medium showed both slant, and butt alkaline without H<sub>2</sub>S production. There was no hemolysis on blood agar culture plate. Further identification was done by VITEK-2 (fully automated identification system) using GN-25 card (bioMerieux, Marcy l'Etoile France). It was identified as *C. testosteroni* with 98% probability. There was no growth in anaerobic blood culture bottle even after 7 days of anaerobic incubation. Urine sample was also sent to the microbiology department for culture and sensitivity. It was processed on Cysteine lactose electrolyte deficient media and similar colony morphology was seen as on MacConkey and blood agar. Further identification was done by VITEK-2 using GN-25 card, and again it was identified as *C. testosteroni* with 97% probability. The isolation of same bacteria from both the clinical samples (blood and urine) confirmed its pathogenic role. The isolate was found to be sensitive to gentamicin, amikacin, imipenem, meropenem, ciprofloxacin, tigecycline, cotrimoxazole, cefuroxime, colistin, and resistant to piperacilline-tazobactam. Therapy was changed to gentamicin 4 mg/kg/dose daily and imipenem 25 mg/kg/dose 8 hourly for 10 days. The patient responded well to antibiotic therapy. Fever was subsided 3<sup>rd</sup> day after starting therapy. Blood sample was taken again for culture, 7 days after completion of antibiotic therapy, and it was negative. Urine sample was also collected with complete aseptic precaution for culture and there was no growth. The patient is doing well during follow-up.

## Discussion

*C. testosteroni* is a widely distributed aerobic Gram-negative bacillus which is rarely seen as a human pathogen. Now a day, this bacterium has been identified in various hospital devices; however, it has not been recognized as a component of the endogenous human microflora. Because this bacterium has been more frequently encountered (till now around 36 cases), we thought that this case would be beneficial for clinicians and laboratory physicians in their diagnosis and clinical approach<sup>[4]</sup> [Table 1]. Majority of the infections are community-acquired rather than nosocomial.<sup>[5,6]</sup> Most of the previously reported patients were immunodeficient due to malignancy, chronic liver disease, and end-stage renal disease taking hemodialysis.<sup>[7,8]</sup> However, in our case, the patient was immunocompetent based on the results of their previous reports. *C. testosteroni* is the most common species among its various species causing human infection and majority of the patients were survived. Till now around 16 cases of *C. testosteroni* bacteremia reported in medical literature, and out of which majority were male but our case

**Table 1: *Comamonas testosteroni* infections in the medical literature**

Age/sex	Diagnosis	Sample	Antibiotic treatment	Result	References
54 years/female	Catheter related infection	Blood	Cefepime, ciprofloxacin	Recovered	Abraham <i>et al.</i>
50 years/male	Cholesteatoma	CSF	Meropenem	Recovered	Arda <i>et al.</i>
31 years/female	Bacteremia	Bone marrow	Kanamycin, tetracycline	Recovered	Atkinson <i>et al.</i>
31 years/male	Appendicitis	Abscess	Cefoxitin	Recovered	Barbaro <i>et al.</i>
24 years/female	IVDA	CSF	Moxalactam and nafcillin	Recovered	Barbaro <i>et al.</i>
59 year/female	Alcoholic cirrhosis	Peritoneal fluid	Cefoxitin	Recovered	Barbaro <i>et al.</i>
11 years/male	Appendicitis	Peritoneal fluid	Ampicillin, clindamycin, tobramycin	Recovered	Barbaro <i>et al.</i>
12 years/female	Appendicitis	Peritoneal fluid	Cefoxitin	Recovered	Barbaro <i>et al.</i>
21 years/female	Appendicitis	Peritoneal fluid	Cefoxitin	Recovered	Barbaro <i>et al.</i>
NB/Stillbirth	Maternal IVDA	Umbilical cord-blood	-	Stillbirth	Barbaro <i>et al.</i>
84 years/female	CHF	Urine	Ampicillin	Recovered	Barbaro <i>et al.</i>
24 years/male	Appendicitis	Peritoneal fluid	Cefoxitin	Recovered	Barbaro <i>et al.</i>
Female/NB	Maternal IVDA	Blood	Ampicillin, amikacin	Dead	Barbaro <i>et al.</i>
49 years/male	Infective endocarditis	Blood and mitral valve	Cefepime, gentamicin	Recovered	Cooper <i>et al.</i>
No data	AIDS	Respiratory secretions	Ceftazidime	Recovered	Franzetti <i>et al.</i>
22 years/male	Appendicitis	Peritoneal fluid, blood	Cefazolin	Recovered	Gul M <i>et al.</i>
75 years/female	Catheter related infection	Blood and CVC	Ceftazidime, gentamicin	Recovered	Le Moal <i>et al.</i>
54 years/male	Cellulitis	Blood	Oxacillin, ciprofloxacin	Recovered	Tsui <i>et al.</i>
73 years/male	Intra-abdominal infection and hepatocellular carcinoma	Blood	Gentamicin, levofloxacin	Recovered	Tsui <i>et al.</i>
82 years/female	Endophthalmitis	Vitreous biopsy	-	Recovered	Reddy <i>et al.</i>
83 years/male	Bacteremia	Blood	Amikacin, piperacillin-tazobactam	Recovered	Katircioglu <i>et al.</i>
63 years/female	Catheter related bloodstream infection in patient on hemodialysis	Blood	Vancomycin, ceftriaxone	Dead	Nseir <i>et al.</i>
10 years/male	Bacteremia	Blood	Ciprofloxacin, amikacin	Recovered	Farshad <i>et al.</i>
19 years/female	Bacteremia	Blood	Ciprofloxacin, vancomycin, imipenem	Recovered	Farshad <i>et al.</i>
51 years/male	Endocarditis	Aortic valve	Ciprofloxacin	Recovered	Arzu Duran <i>et al.</i>
80 years/female	Bacteremia	Blood	Cefazolin and doripenem	Recovered	Orsini Jose <i>et al.</i>
42 years/female	Septic shock	Blood and urine	Ceftazidime and levofloxacin	Recovered	Hyun Jung Kim <i>et al.</i>
65 years/female	Gastroenteritis	Stool	Ciprofloxacin and probiotics	Recovered	Shaika Farooq <i>et al.</i>
46 years/female	Bacteremia	Blood and urine	Gentamicin and imipenem	Recovered	Present case

CVC=Central venous catheter, CSF=Cerebrospinal fluid, CHF=Congestive heart failure, AIDS=Acquired immune deficiency syndrome, IVDA=Intravenous drug abuse, NB=Newborn

was a female patient. Most of the patient responded well to appropriate antimicrobial therapy. *Comamonas* species are usually susceptible to aminoglycosides, fluoroquinolones, carbapenems, piperacillin-tazobactam, trimethoprim-sulfamethoxazole, and cephalosporins.

Until 1987, *C. testosteroni* has only rarely been associated to human infections. In the late 1980s, researchers reported 18 unrelated cases during a research >3 years suggesting that this species might be missed if not carefully identified by clinical microbiology laboratories. Although the pathogenic potential of *C. testosteroni* has not been well studied, it is usually considered a commensal. Because *C. testosteroni* can be isolated from fresh water, it seems likely that wound infection could occur after contact with contaminated water.

Intra-abdominal infections are the most common infections caused by *C. testosteroni*. Bacterial

translocation from gastrointestinal tract seems to play an important role in the pathogenesis caused by *Comamonas* species.<sup>[9]</sup> The high genetic pliability of *Comamonas* species, as well as their inherent ability as an environmental pathogen to survive in various ecological niches, makes it an appropriate candidate to cause mild but persistent infections, particularly in an individual with predisposing conditions. In our case, there was no obvious source of infection. The patient did not have central venous catheters, USG abdomen did not show any mass. The patient did not have any immunological deficiency such as malignancy and diabetes mellitus. HIV status was negative. All these factors might have played a role as predisposing factors. In the present case, the patient belongs to rural area so the source of infection might be the contaminated food and water, although this hypothesis cannot be proven. It is only presumed as there was no proven source of infection.

Polymicrobial bacteremia involving *C. testosteroni* have been reported in association with *Streptococcus parasanguis*, *Streptococcus agalactiae*, *Ralstonia pickettii*, and *Bacteroides fragilis*.<sup>[10]</sup> *C. testosteroni* is able to survive in the hospital environment for a long time, and majority of infections are community acquired. It is anticipated that nosocomial infection related to this bacterium will increase over time as there is increase in long hospital stay with impaired immune system and long-term use of the intravenous catheter. Most of the laboratories are not well equipped to detect such atypical pathogens that can cause human infection in certain clinical settings. Therefore, automated identification system (VITEK-2) can help to identify such rare isolates. In this case report, we have highlighted an unusual cause of bloodstream infection caused by *C. testosteroni*. *Comamonas* species should be kept in mind as a rare cause of bacteremia. The main target to report such an unusual case is to sensitize clinicians and clinical microbiologists for bloodstream infections caused by such unusual pathogens. However, more research should be done to understand the pathogenicity, virulence mechanism, and increased antibiotic resistance of *C. testosteroni*.

### Declaration of patient consent

The authors certify that they have obtained all appropriate patient consent forms. In the form the patient(s) has/have given his/her/their consent for his/her/their images and other clinical information to be reported in the journal. The patients understand that their names and initials will not be published and due efforts will be made to conceal their identity, but anonymity cannot be guaranteed.

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Nil.

### Conflicts of interest

There are no conflicts of interest.

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