

New enterant in the class of uropathogens-*Shewanella algae*

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ABSTRACT

S. algae is considered a rare opportunistic pathogen for humans, frequently involving immune-compromised hosts and are usually part of a polymicrobial infection which may mask its clinical importance. We are reporting first ever case report of urinary tract infection caused by *Shewanella algae*. This is a case of 70 year old diabetic male presented with fever, urgency of micturition, oliguria, swelling over both legs. His midstream urine sample was sent for microbiological examination and it revealed *Shewanella algae*. Patient was treated with antibiotics and he recovered completely after treatment. This case highlights the need to consider *Shewanella* as a potential emerging uropathogen, and utmost microbiological vigilance is required to identify this unusual agent of UTI.

Key words: Diabetic, *Shewanella algae*, Uropathogen

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INTRODUCTION

In 1985, MacDonell and Colwell proposed the new genus *Shewanella*, which was initially composed of three species: *S. putrefaciens*, *S. hanedai* and *S. benthica*.¹ Currently, there are at least 22 species in genus *Shewanella*. Most of them are associated with aquatic and marine habitat.² In 1990, Simidu et al. proposed the name *S. algae* for a tetrodotoxin-producing isolate recovered from red algae.³ Previously, *Shewanella algae* was known as *Alteromonas putrefaciens* or *Achromobacter putrefaciens*.⁴ *Shewanella spp.* has been implicated in skin and soft tissue infections, bacteraemia, biliary tract infections, thoracic empyema, endocarditis, dacryocystitis, intracranial abscess, arthritis, peritonitis, ventilator-associated pneumonia, and ear infections.⁵ In Denmark, 67 ear infections caused by *S. algae* have been published.⁶ We have not come across any case of urinary tract infection caused by *S. algae*.

Two *Shewanella species*, *S. algae* and *S. putrefaciens*, have been found in clinical specimens. Because automated systems are unable to distinguish between two species, a number of infections attributed to *S. putrefaciens* probably correspond to *S. algae*.⁷ *S. algae* is considered

a rare opportunistic pathogen for humans, frequently involving immune-compromised hosts and are usually part of a polymicrobial infection which may mask its clinical importance. We are reporting first ever case report of urinary tract infection caused by *Shewanella algae*.

CASE REPORT

A 70 year old diabetic male was admitted in the Department of Medicine with complaints of fever since 15 days, urgency of micturition, oliguria, swelling over both legs since 4-5 months. He had a past history of being operated for 7mm left renal stones 3 years back. Since then he was having recurrent episodes of urinary tract infections.

Microbiological examination

A 5 ml clean-catch midstream urine sample was collected and was sent to the department of Microbiology for examination.

On Macroscopic examination, it was yellow and turbid urine sample. The sample was inoculated on Cysteine Lactose Electrolyte Deficient (CLED) agar by semi quantitative method and kept for incubation at 37°C for 16-18 hours.

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Microscopic examination showed 10-15 Polymorphonuclear cells/hpf (PMNs) and motile bacilli. On Gram staining there was evidence of Gram negative bacilli and PMNs.

The next day a significant count of 10⁵ Colony Forming Unit (CFU)/ml of mono-microbial non lactose fermenting colony was found on CLED agar.⁸ The colony was subjected to catalase test, oxidase test and standard biochemical tests were applied. Antibiotic susceptibility testing was done by Kirby-Bauer disc diffusion method using discs of standard concentrations provided by Hi - media laboratory (Mumbai, India). Minimum Inhibitory Concentration (MIC) was determined by agar dilution method for ampicillin and ceftriaxone.

RESULTS

Catalase and Oxidase were produced.

Biochemical reactions – (from right to left)

TSI-alkali/acid with H₂S. Indole was not produced. Methyl red and VogesProskauer test were negative. Citrate was utilised. Urease was not produced (Figure 1).

Antibiotic susceptibility pattern

It was found to be sensitive to gentamycin(18mm), ciprofloxacin(27mm), amikacin(23mm), cefepime(21mm), meropenem(32mm), ceftazidime(28mm), ceftriaxone (30mm), aztreonam(26mm), norfloxacin (24mm), nitrofluorantoin(27mm) and was resistant to ampicillin(6mm) and cephalothin(6mm).⁹

MIC was done by agar dilution and it was found resistant to ampicillin (MIC ≥ 32µg/ml) and sensitive to ceftriaxone (MIC ≤ 1µg/ml).

As *Shewanella spp* is only non lactose fermenting, oxidase producing organism giving H₂S on TSI it was subjected to further examination for identification of species. It was inoculated on Blood agar and MacConkey agar for confirmation.

On blood agar-1-2 mm, circular, convex, smooth, beta-haemolytic colonies, with orange tan pigment were seen (Figure 2).

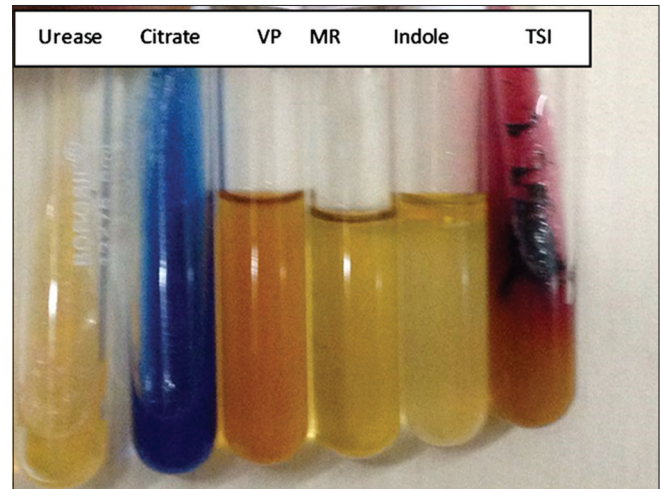


Figure 1: Biochemical properties of *Shewanella* algae

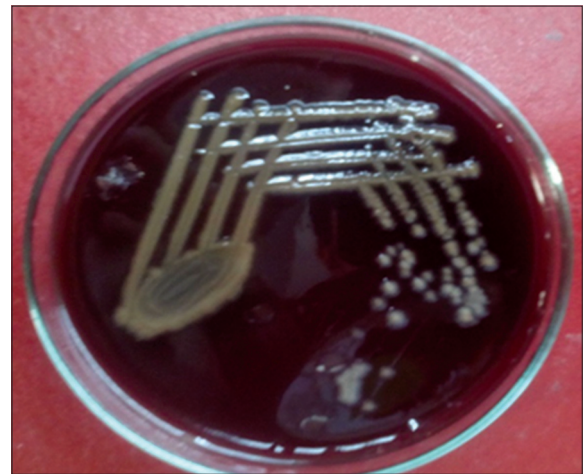


Figure 2: Colonies on blood agar.⁴



Figure 3: Ornithine decarboxylation.

| Table 1: Biochemical tests done for Species level identification of <i>S. algae</i> | |
|---|---|
| Tests done | Result |
| Nitrate reduction | Nitrate was reduced to nitrite |
| Ornithine decarboxylation | Ornithine was decarboxylated (Figure 3) |
| DNase test | DNA was hydrolysed |
| ONPG (O-Nitro Phenol β Galactosidase) | Positive |
| Oxidation-Fermentation test for Glucose and maltose | Glucose and Maltose Oxidised |

The isolate was thermo-tolerant with presence of growth at 42°C and halophilic with growth in presence of 6.5% NaCl.

The biochemical tests done for speciation of *Shewanella* algae are as mentioned in Table 1.

Urine biochemistry examination-

Sugar-trace, albumin-2+.

Blood investigations in biochemistry-

Blood sugar - Post prandial was 208mg% and fasting was 138mg%,

Serum creatinine was 1.6.

Pathological examination-Haemoglobin was 11.2gm%,

White Blood Count -10,700/cmm.

Differential Leucocyte Count- Neutrophils-72%, lymphocytes-25%, monocytes-1%, basophils- 0, eosinophils 2%, Erythrocyte Sedimentation Rate -42mm/l, C-Reactive Protein -23mg/l.

DISCUSSION

Shewanella species are found throughout the world in marine environments and most reported human infections occur in countries with warm climates.²

Shewanella algae and *Shewanella putrefaciens* are the two species of *Shewanella* that are most frequently implicated in human infections. Among *Shewanella* spp. *S. algae* causes 80% human infections.¹⁰

S. algae grows at 42°C and in presence of 6.5% NaCl and this helps to distinguish it from *S. putrefaciens*.⁴ *Alishewanella* is also halophilic but it is non motile and doesnot produce H₂S unlike *S. algae*.⁴ The isolate can be further confirmed to the species level by 16S ribosomal DNA sequencing by using the primers B27F (5'-AGAGTTTGATCCTGGCTCAG-3') and U1492R (5'-GGTTACCTTGTTACGACTT-3') but was not done in our case.¹¹

Patient was admitted for urinary complaints so the organism is most likely to be a community acquired rather than hospital associated pathogen. Patient was treated as a case of recurrent UTI but the clinical signs and raised albumin level with history of diabetes suggest of some pathology like diabetic nephropathy but further investigations like renal ultrasound was not done in this case so we are not sure of any long standing illness in our case. The organism generally resides in marine and aquatic environment so the route of transmission in this case is not known. There have been only four reports of isolation of *Shewanella* from India and these were from patients with infective endocarditis, peritonitis, and chronic obstructive

pulmonary disease respectively.^{12,13} *Shewanella* was isolated in monomicrobialform with a significant bacterial count and clinical picture suggestive of acute urinary tract infection. Patient was treated with a five days course of Norfloxacin 400 mg twice daily. Patient was improving clinically and his symptoms were relieved. A repeat sample was taken on seventh day was sterile. So we believe that *Shewanella* algae was the causative agent in our case.

There are no standard guidelines for treatment of *Shewanella* infection. In present study *S. algae* was found to be susceptible to gentamicin, norfloxacin, amikacin, cefepime, meropenem, ceftazidime, cefotaxime, aztreonam, nitrofurantoin and resistant to ampicillin and cephalothin. According to some textbooks resistance is found to ampicillin and cephalothin.⁴ Previously published studies reported that *S. algae* is susceptible to aminoglycosides, carbapenems, erythromycin, and quinolones, with variable susceptibility to penicillin and cephalosporins. *S. algae* is characteristically susceptible to aminoglycosides, carbapenems, erythromycin and quinolones, but resistant to penicillin. Susceptibility to ampicillin and cephalosporins is variable, with more isolates being susceptible to third- and fourth- than to first- and second-generation cephalosporins.¹⁴ However, rapid development of resistance to imipenem and piperacillin-tazobactam has been reported.¹⁴

To conclude, this case highlights the need to consider *Shewanella* as a potential emerging uropathogen, and utmost microbiological vigilance is required to identify this unusual agent of UTI.

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Authors Contribution:

OK - Isolation of Pathogen, guidance in identification, revised the manuscript; **PS, SP & AJ** - history taking, performed Lab tests, drafted the manuscript; **SN** - Reviewed & finalised the manuscript.

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