Pseudomonas mendocina Meningitis in a Postoperative Patient: A Case Report and Review of Literature

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ABSTRACT

Microbiology Section

Pseudomonas mendocina is a Gram negative bacillus belonging to the Pseudomonadaceae family. It was first isolated in 1970 from soil and water in the Mendoza region of Argentina. Although it rarely causes infection, 21 cases of infections ranging from infective endocarditis, bacteraemia, meningitis, soft tissue infections, and urinary tract infections have been reported worldwide, mostly among patients with underlying co-morbidities. This report is the first case of pyogenic meningitis due to *Pseudomonas mendocina* reported from a 1600-bed teaching hospital in Northern India and the second case of *Pseudomonas mendocina* infection from the Indian subcontinent. The presented case was of a 31-year-old female who was diagnosed with pituitary macroadenoma on Magnetic Resonance Imaging (MRI). She underwent endoscopic endonasal trans-sphenoidal excision of the tumour after two months of diagnosing the tumour, following which she developed meningitis. On two consecutive aerobic bacterial cultures and identification by Matrix-Assisted Laser Desorption/Ionisation-Time of Flight-Mass Spectrometry (MALDI-TOF-MS), growth of *Pseudomonas mendocina* and Amoxicillin-clavulanic acid which alleviated her symptoms of meningitis and she was discharged after 27 days of hospital stay.

Keywords: Bacterial meningitis, Cushing's disease, Immunocompromised patients, Opportunistic infections, Pituitary macroadenoma

CASE REPORT

A 31-year-old female, presented to the endocrinology Outpatient Department (OPD) with chief complaints of headache and temporal vision loss for the past 20 days. She was advised to undergo an MRI brain. Her MRI was suggestive of pituitary macroadenoma, as represented by [Table/Fig-1]. Other clinical features like raised blood pressure of 142/98 mmHg, raised blood sugar levels of 346 mg/dL and osteoporotic changes in bones were suggestive of Cushing's disease with no family history of the disease. She was admitted to the endocrinology department for treatment of diabetes insipidus, osteoporosis and was started on cortisol to manage her Cushing's disease and sodium levels were continuously monitored. Due to temporal heminopsia, she was advised for decompression of the optic nerve and further referred to the Neurosurgery department for evaluating the chances of surgically excising the pituitary tumour. Maintaining aseptic precautions, endoscopic endonasal trans-sphenoidal excision of the tumour was performed after two months of diagnosing the tumour. After the operation, patient was shifted in intubated and unreversed state to the Intensive Care Unit (ICU). Patient was extubated the next



Pituitary macroadenoma (shown by yellow coloured arrow in the image) in the patient before surgery.

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day and the postoperative Computed Tomography (CT) of the head showed a well formed surgical corridor with near total excision of tumour showing evidence of temporal horn pneumocephalous with no evidence of haematoma or infarct, as seen in [Table/Fig-2]. Postoperatively, her requirement for antihypertensive drugs ceased. On her fourth day postsurgery, the patient developed headache and fever for which routine Cerebrospinal Fluid (CSF) examination was performed as shown in [Table/Fig-3], which was suggestive of pyogenic meningitis.



[Table/Fig-2]: Computed Tomography (CT) image of brain showing well formed surgical corridor (yellow arrow) with near total excision of tumour showing evidence of temporal horn pneumocephalous with no evidence of haematoma or infarct postsurgery.

Routine parameters of CSF for detection of infection	Normal parameters	Parameters of the CSF in case of the patient					
Total leukocyte count (per cubic millimeter)	0 to 5	102					
CSF glucose (mg/dL)	50 to 80	30					
CSF protein (mg/dL)	15 to 40	128					
CSF lactate (mmol/ liter)	1-3	9					
Microbial examination	No microorganism	Gram negative bacilli seen					
[Table/Fig-3]: Routine Cerebrospinal Fluid (CSF) examination suggestive of pyogenic meningitis.							

The CSF sample was sent to Bacteriology section of the Department of Microbiology at a tertiary care centre where first a

wet preparation and India ink preparation was performed which showed moderate pus cells, few red blood cells, few motile bacilli and no capsulated microorganisms and simultaneously a Grams stained smear was prepared which showed few pus cells and few Gram negative bacilli, as shown in [Table/ Fig-4]. The sample was also inoculated on MacConkey agar and blood agar followed by Robertsons' Cooked Meat (RCM) broth. After 16-18 hours of incubation at 37°C, growth of non lactose fermenting colonies were observed on MacConkey agar, while the growth on blood agar was non haemolytic and pale colonies were observed on nutrient agar, as shown in [Table/ Fig-5]. The microorganism was identified as Pseudomonas mendocina by MALDI-TOF-MS (Bruker Daltonics, Germany) assay. To confirm that the microorganism was not a mere environmental contaminant, authors requested the clinicians for a repeat sample and it revealed the growth of the same microorganism. A pair of BACTEC blood culture bottles from the patient was also sent on the same day as the repeat CSF sample, but did not flag positive. AST of the isolate using Kirby Bauer disc diffusion method as per the Clinical and Laboratory Standard Institute (CLSI) 2019 guidelines was performed [1]. The isolate was resistant to amikacin, aztreonam, ceftazidime, ciprofloxacin, cefoperazone- sulbactum, imipenem, meropenem and piperacillin- tazobactam and was only found susceptible to colistin with Minimum Inhibitory Concentration (MIC) of 1 µg/mL. As the clinicians did not want to start an antibiotic of last resort in this case they the patient was started on piperacillin-tazobactam 4.5 gram and amoxicillin- clavulinic acid 625 mg TDS and was relieved of fever and headache after 10 days of treatment. The use of a resistant antibiotic could obtain response which can be justified by the 90:60 rule which states that infections due to resistant isolates respond approximately 60% of the time. The 90:60 rule worked in their favour and on discharge she was continued on amoxicillin- clavulinic acid 625 mg TDS, cortisol 100 µg and wyslone (steroid) 5 mg for 14 days. The patient had



[Table/Fig-4]: Grams stained smear showing few pus cells and few Gram negative bacilli (yellow arrow) observed under 100x magnification of a compound microscope.

improved completely at the time of discharge and was advised for follow-up in the neurosurgery OPD after six weeks or in case of altered sensorium, high grade fever and seizure or CSF leak. The patient did not return for follow-up and so her progress could not be recorded any further.

DISCUSSION

Pseudomonas mendocina, belonging to the Pseudomonadaceae family is a gram negative, aerobic and non sporing bacillus. It was originally identified from soil and water of Mendoza region in Argentina in 1970; hence it was named after the region of origin [2]. The microorganism belongs to the Pseudomonas putida Group-II and was not recognised to be the cause of infections among humans [3]. The case that deemed this rare organism as a pathogen was infective endocarditis in patient who had undergone aortic valve replacement surgery in Mendoza, Argentina in the year 1992 [4]. Following which, many cases of infection by this rare pathogen in immunocompromised patients have been reported, which include 21 known cases of infections which have been documented from Asia, Europe, Middle-east, North and South America, as demonstrated in [Table/Fig-6] [3-19]. Pseudomonas mendocina is a known cause of hospital acquired and opportunistic infections among the immunocompromised patients [20]. Although the microorganism is seldom known to cause infection, its appearance in clinical sample from a patient can account for nosocomial encounter with this pathogen in the duration of prolonged hospital stay.

Seven cases of infections have been reported from South-East Asia by Howe TS et al., Chiu LQ and Wang W, Huang CR et al., and Gupta V et al., of which five cases were immunocompromised while two cases were reported in immunocompetent patients [10,11,15,18]. *Pseudomonas mendocina* wound infection in a diabetic and asthmatic farmer, who was intermittently on steroids, was first recognised by Gupta V et al., in India [18]. Four cases of pyogenic meningitis reported from Taiwan have been known to be caused by this pathogen in 2018 [15].



[Table/Fig-5]: Nutrient agar, MacConkey agar and Blood agar showing colonies of Pseudomonas mendocina.

Publication year	Author	Country of publication	Age	Sex	Co-morbidities	Infection
1992	Aragone MR et al., [4]	Argentina	63	Male	Diabetes mellitus type 2, aortic valve replacement, poliomyelitis	Infective endocarditis
2001	Johansen HK et al., [5]	Denmark	28	Female	Situs inversus, double-outlet right ventricle, Ventricular Septal Defect (VSD), pulmonary stenosis, multiple cardiovascular surgeries	Infective endocarditis
2005	Chi CY et al., [6]	Taiwan	65	Male	Alcoholic hepatitis, chronic renal disease	Spondylodiscitis
2007	Mert A et al., [7]	Turkey	36	Male	Mental retardation	Infective endocarditis
2011	Suel P et al., [8]	France	79	Female	Atrial fibrillation, transient ischemic attack, hypertension	Infective endocarditis
2011	Nseir W et al., [9]	Israel	31	Male	Healthy	Bacteraemia
2013	Howe TS et al., [10]	Singapore	86	Female	Vertebral compression fractures, tibial plateau stress fracture	Osteomyelitis
2013	Chiu LQ and Wang W [11]	Singapore	34	Male	Healthy	Septic arthritis
2016	Rapsinski TM et al., [12]	United States	57	Male	Gout, chronic alcohol use	Infective endocarditis

2017	Jerónimo TM et al., [13]	Portugal	22	Male	Chronic kidney disease, peritoneal dialysis	Peritonitis	
2018	Almuzara M et al., [14]	Argentina	56	Male	Alcohol use disorder, vascular insufficiency	Burn wound infection	
2018	Almuzara M et al., [14]	Argentina	36	Male	Alcohol use disorder	Burn wound infection	
2018	Huang CR et al., [15]	Taiwan	55	Male	Diabetes mellitus type 2, buccal cancer, community-acquired infection	Meningitis	
2018	Huang CR et al., [15]	Taiwan	66	Female	Spontaneous intracerebral haemorrhage, external ventricular drainage	Meningitis	
2018	Huang CR et al., [15]	Taiwan	79	Male	Chronic obstructive pulmonary disease, respiratory failure, nosocomial infection	Meningitis	
2018	Huang CR et al., [15]	Taiwan	78	Female	Healthy	Meningitis	
2019	Gani M et al., [3]	United States	63	Male	Resistant HIV/AIDS	Bacteraemia	
2020	Goldberg ME et al., [16]	United States	72	Male	End-stage renal disease, immunoglobulin A (IgA) nephropathy, atrial fibrillation, heart failure with reduced ejection fraction, obesity, chronic venous stasis	Bacteraemia	
2021	Ezeokoli EU et al., [17]	United States	81	Male	Coronary artery disease, atrial fibrillation, heart failure, chronic kidney disease, diabetes mellitus type 2, cerebrovascular accident	Bacteraemia	
2021	Gupta V et al., [18]	India	53	Male	Diabetes mellitus type 2, asthma	Leg wound infection	
2022	Vo T et., al [19]	United States	83	Male	Diabetes mellitus type 2, hypertension, coronary artery disease, prostate cancer, COVID-19 pneumonia	Urinary tract infection	
2023	Present case	India	31	Female	Pituitary macroadenoma with secondary Cushing disease	Meningitis	
[Table/Fig-6]: Review of literature on 21 cases of infections caused by Pseudomonas mendocina till date (N=22) [3-19].							

Authors have recognised the first case of *Pseudomonas mendocina* as a causative agent of pyogenic bacterial meningitis at a Tertiary care centre in Northern India. Earlier cases of bacterial meningitis have been reported from Taiwan in the year 2018 by Huang CR et al., [15]. The preponderance of none of the genders was observed in the studies conducted by them as out of the four patients, 50% were male. All the patients included in the reports by Huang CR et al., were above 50 years of age and had underlying immunocompromised conditions that rendered them easily susceptible to opportunistic infections by rare pathogens, in comparison to index case 31-year-old patient suffering from pituitary macroadenoma [15].

The patient in present case report was admitted for a period of 27 days and during the course of hospital stay after the four days postoperatively she was diagnosed of pyogenic meningitis caused by this rare isolate. Due to her immunocompromised status and underlying co-morbidities including continuous use of cortisol for Cushing's disease, the host was vulnerable to acquire opportunistic infections from the hospital environment as a rare pathogen like *Pseudomonas mendocina* which is an inhabitant of the water and soil [2]. Another cause of infection could be the poor compliance of the hospital staff to infection control measures and hand washing protocol, but the exact source of infection was unidentifiable.

Being a rare pathogen, there was faint knowledge about its antibiotic susceptibility pattern. As *Pseudomonas mendocina* belongs to the *Pseudomonas putida* group (Group-II), which is susceptible to most routine antibiotics, on the contrary, the isolate was rather resistant to most antibiotics [12]. Thus, identification of resistant isolates of *Pseudomonas mendocina* and administration of appropriate antibiotics and their dose adjustment can reduce morbidity among postoperative patients due to meningitis and expedite their recovery. This was the second case of *Pseudomonas mendocina* and the first case of *Pseudomonas mendocina* mendocina mendocina mendocina

CONCLUSION(S)

Pseudomonas mendocina is a rare pathogen known to cause infective endocarditis, bacteraemia, meningitis, and wound ulcers. This is the fifth reported case of pyogenic meningitis in the world and the first case from India. This case demonstrates the second *Pseudomonas mendocina* related infection in the Indian

subcontinent, and identifies the patient's immunocompromised state as being responsible for the acquisition of opportunistic infections caused by uncommon pathogens.

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