A Case of Lower Respiratory Tract Infection with Canine-associated *Pasteurella canis* in a Patient with Chronic Obstructive Pulmonary Disease

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ABSTRACT

Microbiology Section

This is the report of lower respiratory tract infection with *Pasteurella canis* in a chronic obstructive pulmonary disease (COPD) patient with history of casual exposure to cats. *Pasteurella* species are part of the oral and gastrointestinal flora in the canine animals. These organisms are usually implicated in wound infection following animal bites, but can also be associated with a variety of infections including respiratory tract infections.

CASE REPORT

A 70-year-old male, hotel employee by occupation, known case of Chronic obstructive pulmonary disease (COPD) and ischaemic heart disease (IHD) presented to our hospital with a history of cough with purulent expectoration, low grade fever and worsening breathlessness of seven days duration. Patient had history of recurrent exacerbations of COPD caused by *Pseudomonas* spp. six months back. Patient was an active smoker and gave a history of casual exposure to domestic cats.

On examination, patient was conscious, afebrile, tachypneic (respiratory rate of 22/minute), mildly hypoxic (oxygen saturation on room air of 88% by pulse oximetry) and haemodynamically stable. Respiratory system examination revealed a barrel shaped chest and bilaterally diminished breath sounds with diffused polyphonic wheeze on auscultation. Routine blood investigations like haemogram, ESR, fasting blood glucose, renal function, serum electrolytes and liver function were all within normal limits. Arterial blood gas analysis was suggestive of mild Type I respiratory failure (pH= 7.36, PaCO₂ = 36 mmHg, PaO₂ = 59.6 mmHg, PaHCO₂ = 21 mmHg). Chest radiograph showed changes of hyperinflation, unfolding of aorta and no evidence of lung parenchymal abnormalities [Table/ Fig-1]. Spirometry was suggestive of severe obstructive impairment with no significant bronchodilator reversibility. Sputum was sent for gram stain, bacteriological culture and sensitivity testing. Patient was treated with low flow oxygen, ceftriaxone 1 gram intravenously BID, hydrocortisone and salbutamol + ipratropium nebulisations. Response to initial therapy at the end of 48 hours was poor.

Gram stain smear of the sputum revealed numerous polymorphonuclear leucocytes with gram negative coccobacilli and it was decided to wait for the culture report before modifying the empiric antibiotic.

The sample was cultured on blood agar, chocolate agar and MacConkey's agar plates and incubated at 37°C for 24 hours. Blood agar plates showed non haemolytic small dew drop colonies and chocolate agar plates showed small grey coloured colonies and the smear from the colonies showed the presence of gram negative coccobacilli [Table/Fig-2,3]. There was no growth on MacConkey's agar plate. The isolate was catalase and oxidase test positive. The isolate was further identified as *Pasteurella canis* by Vitek 2 system (Bio-Mérieux, Co., Ltd.). Antibiotic susceptibility testing was done by modified Kirby-Bauer disk diffusion technique. The organism

Keywords: Canine animals, Doxycycline, Vitek 2 system



[Table/Fig-1]: Chest radiograph PA view showing hyper-inflated lung fields and an unfolded aorta [Table/Fig-2]: Culture on Chocolate agar plate showing smooth grev colonies of *P.canis*



[Table/Fig-3]: Gram smear from the growth showing gram negative cocco bacilli [Table/Fig-4]: Antibiotic Susceptibility testing: plate showing zone of inhibition to Levofloxacin, Erythromycin, Gentamycin, Tetracycline

was sensitive to ciprofloxacin, amoxicillin-clavulanic acid, penicillin, gentamicin, clindamycin, levofloxacin, erythromycin, doxycycline and trimethoprim-sulfamethoxazole ([Table/Fig-4], interpretation with *Haemophilus influenzae* standards).

Taking into consideration the antibiotic sensitivity, parenteral ceftriaxone was stopped and replaced by oral doxycycline along with other supportive care. The patient showed gradual improvement and was discharged on oral doxycycline for two weeks along with inhaled bronchodilators. On follow up at two weeks, he was asymptomatic and a repeat sputum culture revealed no significant bacterial growth.

DISCUSSION

Pasteurella canis is a gram-negative, non-motile coccobacillus or short rods belonging to the Pasteurellaceae family [1]. First referred to as "*Micrococcus gallicidus*", the generic name was redesignated as "*Pasteurella*" in 1887 by Trevisan to commemorate the work of Pasteur on these bacteria. Like most species of *Pasteurella*, *P.*

canis is oxidase and catalase test positive. It includes two biotypes: biovar 1 is originated from canine, whereas biovar 2 is originated from bovine animals. The two biotypes are distinguishable by the indole test: biovar 1 is indole positive whereas biovar 2 is indole negative [1].

P. canis are a part of the normal oropharyngeal flora of many animals including healthy dogs and cats. In humans, they are known to cause zoonotic infections. Human pasteurellosis most often results in skin or soft tissue infections after an animal bite. *P. multocida* is most commonly isolated in human infections but there have been reports of other species such as *P. canis* and *P. dogmatis* being involved [2]. *P. canis* is usually transmitted to human through animal bites, licks. Dog bites are most commonly implicated followed by cat bites. Exceptionally, some patients develop infections after other animal exposure and in some infection may occur even in the absence of an animal contact [3].

After soft tissue and wound infections, the respiratory tract is the second most common site for *Pasteurella* infection. Most patients with pulmonary infection due to *Pasteurella* are elderly with other pre-existing chronic lung diseases like COPD, bronchiectasis, or malignancy. The list of 'pulmonary pasteurellosis' includes tracheobronchitis, pneumonia, lung abscess and empyema [4]. *P.canis* causing bacteremia, peritoneal dialysis-related peritonitis, ocular infections including conjunctivitis outbreaks, osteomyelitis, cutaneous abscess and septic arthritis in the immunocompromised patients has been reported in the literature as well [5-9].

Pasteurella spp. is known to be susceptible to Penicillin G, amoxicillin-clavulanate, piperacillin, fluoroquinolones (levofloxacin, moxifloxacin), newer generation cephalosporins (ceftriaxone, cefixime, cefpodoxime), doxycycline and carbapenems. Treatment failures have been reported with the use of oral macrolides (e.g. erythromycin), oxacillin, dicloxacillin, first generation cephalosporins and clindamycin which should therefore be avoided [10].

Review of literature did not reveal any previous reports of *P.canis* being implicated as a co-pathogen in COPD exacerbations, although the organism itself finds mention as a causative agent in a multitude of other system disease usually against a background of intimate animal contact or trauma. In our patient, since there was only a casual contact with cats and no history of a scratch or a bite from the animal, we assume that he would have been exposed to secretions of his pet animal through inhalation of contaminated aerosol. The isolation of *P. canis* in the sputum of an elderly patient admitted with a COPD exacerbation and the fact that he had only an insignificant history of feline contact prompted us to report this case.

Kim *et al.*, have also reported a case of respiratory tract infection caused by *P. canis* in a COPD patient (poodle owner). This bacteria is found in the oral secretions of canine animals and it can colonize and infect the respiratory tract in patients with lung disease. The patient was started on doxycycline and the symptoms improved. The presentation of this case is similar to our case [11].

T Akahane et al., have reported dual infection with *Pasteurella* dagmatis and *P.canis* in dog bite wound infection in a 25-year-old

female [12]. The other infections caused by *Pasteurella* species reported in literature include cellulitis, subcutaneous abscesses following dog and cat bite, endocarditis following a cat-bite, vertebral osteomyelitis, spondylodiscitis in a diabetic patient [13,14]. Moreover, first case of association of *P.canis*, with bacteremia in a cirrhotic patient with open leg was reported by Albert et al., [15].

However to the best of our knowledge, this is the first case of exacerbation of COPD with *Pasteurella* species co-infection to be reported from this region.

CONCLUSION

Obtaining a detailed history of animal exposure in COPD patients is of paramount importance for the diagnosis of respiratory tract infection caused by *Pasteurella* spp. Elderly patients with COPD need to avoid close contact with pet animals as this could be a potential risk factor for pneumonia caused by *P. canis*.

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