Nocturnal Hypoxemia, an Unusual Presentation of Pulmonary Endometriosis: A Case Report

Andre B. Zanella\textsuperscript{a}, Pedro P. Oliveira\textsuperscript{a}, Renato Seligman\textsuperscript{a,b}, Beatriz G. S. Seligman\textsuperscript{a}, Tania W. Furlanetto\textsuperscript{b,c}

Abstract

Thoracic endometriosis syndrome is a rare condition with four main presentations: pneumothorax, hemothorax, hemoptysis, and lung nodules. We describe an incidental diagnosis of pulmonary endometriosis in a patient admitted with acute pyelonephritis, and developed acute hypoxemia and a productive cough of blood tinted sputum, on the third day of hospitalization. Her thoracic x-ray showed diffuse pulmonary infiltrates, and a small pleural effusion. Chest computerized tomography scan had extensive areas of ground glass opacifications scattered predominantly in the medulla of both lungs. Bronchoalveolar lavage cytology showed small groups of cells in arrays compatible with endometrial cells, which confirmed the diagnosis of pulmonary endometriosis. On the seventh day of hospitalization, after the end of her menstrual bleeding, oxygen supplementation was discontinued, and hypoxemia did not recur. A chest x-ray at the ninth day of hospitalization was normal. Pulmonary endometriosis should be considered in a patient who has unexplained episodes of nocturnal hypoxemia associated with hemoptysis, especially if there is periodicity.

Keywords: Endometriosis; Pulmonary function; Hypoxemia; Nocturnal

Introduction

Endometriosis is defined as the presence of endometrial-like glands outside the uterine cavity, usually ovaries, cul-de-sac, broad and uterosacral ligaments, and, less commonly, in the abdomen, thorax, brain and skin [1]. Thoracic endometriosis syndrome (TES) is a rare condition with four main presentations: pneumothorax, the most common, hemothorax, hemoptysis, and lung nodules. The diagnosis is usually established clinically when women have recurrent episodes of chest pain, dyspnea, and/or hemoptysis during menstruation. It is commonly associated with pelvic endometriosis [2-4].

Our aim is to describe an incidental diagnosis of pulmonary endometriosis in a 38 year-old patient who was admitted to treat acute pyelonephritis, and developed acute hypoxemia at night.

Case Report

A caucasian 38-year-old woman was admitted to hospital with a chief complaint of fever within the last 24 hours, a left sided lower back pain for 3 days, dysuria and nausea. She presented a previous episode of urinary tract infection one month ago and had a previous medical history of hypertension treated with ACE inhibitors since her second pregnan-
She had no other complaints and denied other previous medical problems. Left lumbar fist percussion was positive; otherwise her physical examination was normal. A urine sample had increased leucocytes, and erythrocytes. Blood hemoglobin was 10.6 g/dL (normal range (NR) 12.0 - 16.0 g/dL), blood leucocytes were 9,180 cells per mL (NR: 4,000 - 10,000 cells/mL), and serum creatinine level was 0.78 mg/dL (NR: 0.5 - 1.2 mg/dL). Urine grew Escherichia coli, and the ultrasound of her urinary tract had no abnormalities. Intravenous cefuroxime was initiated with resolution of symptoms. On the third day of hospitalization at 4:30 AM, during routine vital signs evaluation, oxygen saturation assessed by pulse oximetry was 70%, and the respiratory rate (RR) was 26 respirations per minute. Oxygen was supplemented by nasal catheter, with improvement of oxygen saturation, and was discontinued at daytime. At 23:15 PM, her oxygen saturation assessed by pulse oximetry was 84%. She was disoriented and complained of not feeling well, especially when lying down, with a productive cough of blood tinted sputum. She presented with tachypnea but denied dyspnea or chest pain. Arterial blood gases showed pH: 7.44 (NR 7.35 - 7.45); pCO₂: 32.3 mmHg (NR 35 - 45 mmHg); HCO₃: 21.6 mEq/L (NR 22 - 26 mEq/L); pO₂: 48.6 mmHg (NR 80 - 100 mmHg), and O₂ saturation: 86.7% (NR 95-100%). This was improved with oxygen supplementation by Venturi mask, and arterial blood gases were pH: 7.44; pCO₂: 36.0 mmHg; HCO₃: 24.0 mEq/L; pO₂: 83.4 mmHg, and O₂ saturation: 97.9%. Her thoracic x-ray showed diffuse pulmonary infiltrates, and a small pleural effusion (Fig.1). The next morning, she informed that her period had begun one day ago, and recalled having epistaxis during her menstrual periods since menarche. She also recalled feeling unwell at night during her menstrual cycles, which were irregular, intensely painful, and had increased blood flow. On the sixth day of hospitalization, a chest CAT scan had no evidence of pulmonary thromboembolism, and showed extensive areas of ground glass opacifications scattered predominantly in the medulla of both lungs (Fig. 2). On the seventh day of hospitalization, after the end of her menstrual bleeding, oxygen supplementation was discontinued, and hypoxemia did not recur. Upper airway endoscopic evaluation was performed and no abnormalities were found. A chest x-ray at the ninth day of hospitalization was normal (Fig. 3). The following day, a bronchoscopy with bronchoalveolar lavage was performed. The bronchial tree was macroscopically normal, and the bronchoalveolar fluid was negative for bacterial infection, tuberculosis, and fungus, and had macrophages (85%), lymphocytes (10%), neutrophils (3%), eosinophils (1%), and basophils (1%). There were macrophages laden with hemosiderin. Peripheral lung and bronchial wall biopsy were normal. Cytology demonstrated small groups of cells in arrays compatible with endometrial cells, which confirmed the diagnosis of pulmonary endometriosis.

Discussion

A woman, with a long history of epistaxis and feeling unwell at night during her period, was hospitalized for pyelonephritis. Hypoxemia during the night led to the diagnosis of pulmonary endometriosis. Although the patient had severe hypoxemia, she did not ask for help, probably because she was asleep when it occurred, so hypoxemia was detected on routine evaluation. It would be expected that hypoxemia would have triggered awakening from sleep, nevertheless, severe oxygen desaturation does not always causes arousal from sleep [5, 6].

Once the abnormality in her blood oxygen levels followed the recumbent position at night, the source of bleeding probably was located in the low areas of her lungs, which allowed blood to spill to non-bleeding areas of the lungs, when recumbent. There was no evidence of hypoventilation, since
the patient had tachypnea and hypocapnia. The increased 
A-an oxygen gradient suggested several mechanisms for hy-
poxemia. Infections, pulmonary thromboembolism, pulmo-
nary congestion, and pulmonary interstitial metastasis were 
excluded by the clinical evolution. The cyclical character of 
her symptoms associated to menstrual flow, and blood tined 
sputum, led to the clinical suspicion of pulmonary endome-
triosis.

Interestingly, the manifestations of TES, except for 
nodules, occurred more commonly in the right hemitho-
rax, clearly an unexplained phenomenon that continues to 
elude all investigators [2, 3]. Diagnosis is usually delayed 
until several episodes have occurred because of failure to as-
sociate the patient’s symptoms with menses [7]. Computed 
tomography scan, unless obtained during menses, may be 
unremarkable [7]. Although histopathological detection of 
thoracic endometrial tissue is generally not required, diag-
nosis was established in some cases by pleural fluid cytology, 
needle biopsy of lung masses, and cytological study of 
bronchoscopic fluid [8-10]. On presentation, patients should 
be managed acutely as other patients with pneumothorax, 
hemothorax or hemoptysis. Once the patient has been sta-
bilized, hormonal manipulation is generally recommended 
as the first line therapy, but recurrence rates are high. When 
hormonal therapy fails or is not tolerated, surgical therapies 
should be considered [7].

**Conclusion**

Pulmonary endometriosis should be considered in a patient who has unexplained episodes of nocturnal hypoxemia associated with hemoptysis, especially if there is periodicity.

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**Conflict Interests**

The authors declare that they have no competing interests.

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