CHRONIC PULMONARY HISTOPLASOMOSIS IN BRAZIL: REPORT OF TWO CASES WITH CAVITATION DIAGNOSED BY TRANSTHORACIC NEEDLE BIOPSY

Luiz Carlos SEVERO(1), Carlos Fernando Carvalho RIZZON(2), Eliane Wurdig ROESCH(3), Flávio de Mattos OLIVEIRA(4) & Nelson da Silva PORTO(5)

SUMMARY

Two cases of Chronic Pulmonary Histoplasmosis are reported and other eleven cases, collected from Brazilian literature, are commented. After being clinically cured, one of our patients presented an *Aspergillus* fungus ball inside a cavitation in the wall of which *H. capsulatum* was disclosed. Comments are also done on the diagnosis of the mycosis.

**KEYWORDS:** Histoplasmosis; *Histoplasma capsulatum*; Lung; Cavitation; *Aspergillus fumigatus*; Fungus ball; Transthoracic biopsy.

INTRODUCTION

Chronic Pulmonary Histoplasmosis results from exogenous primary infection or reinfection with the propagules of *Histoplasma capsulatum* var. *capsulatum* of an adult patient presenting a lung structural defect. The most frequent predisposing factor has been bullous or centrilobular emphysema, commonly in heavy smokers. Frequently signs and symptoms of chronic obstructive bronchopulmonary disease are associated. The lesions are limited to the lungs. Early and later on clinical and radiological features are similar to those of chronic pulmonary tuberculosis(1,6,10).

Only eleven cases of chronic pulmonary histoplasmosis have been reported in Brazil. Two new cases will be added in this report. In one of the patients an *Aspergillus* fungus ball complicated the disease.

CASE REPORTS

Case 1 – A 62 year-old white man living in Santiago (RS) was admitted on Oct. 17, 1988 with a 7-months history of cough, purulent expectoration, hemoptysis, fever, anorexia, dyspnea and weight loss. The patient was a heavy smoker (30 cigarettes daily for 50 years) and a moderate drinker. His medical history was significant for a partial gastrectomy 22 years before. On physical examination diffuse crackles were heard in both lungs; abdominal examination revealed a median supraumbilical scar of abdominal surgery; and liver edge was palpable 3 cm below the costal margin. A chest X-ray revealed emphysema, scattered nodules in both lungs, small cavities in the apical posterior segment of the left upper lobe, and a cavitation in the right upper lobe, with adjacent pleural thickening (Fig. 1). Tuberculin skin test was negative. No acid-fast bacilli were observed in smeared sputum and in the sediment of the bronchoalveolar lavage; but, small, oval, budding yeast cells and pseudohyphae were disclosed in Grocott stained smears of both these specimens (Fig. 2). A transthoracic cutting needle biopsy was then, performed; in H&E stained sections a fibrotic margin enclosing a palisade of epithelioid cells surrounding a necrotic focus, containing leukocytes, were seen; budding oval yeast cells are disclosed within the necrotic focus by Grocott stain. *H. capsulatum* var. *capsulatum* and *Candida albicans* were isolated from the sputum and the sediment of bronchoalveolar lavage planted on BHI and Sabouraud, incubated, respectively 35°C and 25°C; but, only *H. capsulatum* grew up in the cultures from biopsied tissue. The patient received ketoconazole, 200 mg twice daily, for three weeks, and, was discharged on 200 mg daily.

Two months later he continued treatment with ketoconazole, but his respiratory condition did not improve completely. The chest X-ray (Jan. 09, 1989) of the patient revealed partial regression of the cavitory lesion in the right upper lobe.

Seven months later (May 29, 1989) the patient was readmitted complaining of hemoptysis, weight loss and dyspnea. Physical examination was normal except for decreased
breath sounds in the right upper lung field. The chest roentgenogram revealed markedly regression of pleural thickening and in the thickness of the walls of cavities in both upper lobes. In addition to ketoconazole, he received amphotericin (0.5 mg/kg/day) for 40 days. The patient improved and was discharged on ketoconazole (200 mg twice daily).

Two and half years later (Nov. 25, 1991) the patient was again hospitalized complaining of frequent and severe hemoptysis, productive cough, fever (38°C), dyspnea, and weight loss. Chest X-ray disclosed apical opacities in both lungs and a round mass within a cavity in the upper left lobe (Fig. 3). An immunodiffusion test revealed precipitin bands against A. fumigatus aspergillin. A lobectomy was performed. At cut section of the resected lobe a thick walled cavity, measuring 6 cm in diameter, limited by smooth irregular wall was observed; brownish friable masses, weighing 5 g lied inside the cavity; fibrosis, anthracosis, nodules and micronodules were also seen in the adjacent lung parenchyma. Microscopic examination of histological sections of the nodules and micronodules, stained by H&E, revealed necrotizing tuberculoid granulomas surrounded by a thin layer of hyalinated tissue; oval budding yeast cells, compatible with H. capsulatum were disclosed in the center of the necrotic foci in Grocott stained sections. Histological sections of the masses removed from the cavity revealed that they consisted of a tangled network of hyaline septate hyphae, 3 μm in width. Small fragments of these masses were inoculated on to Sabouraud dextrose agar slants and incubated at both room temperature and at 37°C; luxurious growth of A. fumigatus were obtained.

Post operative course: after surgery the patient’s condition deteriorated (pleural bacterial empyema) and he died.

Case 2 – This patient was a 64 year-old white man living in Ituqui (RS), admitted to our hospital on February, 1990. Since four months ago, because he presented cough, mucopurulent sputum, fever, weight loss, night sweats, and dyspnea he was treated for a presumed tuberculosis, in spite of negative tuberculin skin test and negative results of microscopic examination of sputum smears for acid-fast bacilli. The patient was a heavy smoker (40 cigarettes daily) and moderate drinker for 48 years. Physical examination revealed a tall, slim man with pectus excavatum. Diffuse crackles were heard in both lungs. A chest roentgenogram showed emphysema, cavitary lesions in
both upper lobes and bronchogenic spread to lower lobes (Fig. 4). Multiple sputum smears revealed no acid-fast bacilli. A transsthoracic needle biopsy was performed; small oval yeast-like cells were disclosed in Grocott stained smear (Fig. 5). The isolates obtained on Sabouraud chloramphenicol medium incubated at room temperature were identified as *H. capsulatum* var. *capsulatum* by its dimorphic presentation in subcultures on BHI incubated at 37°C. Immunodiffusion test revealed M band. The patient improved on ketoconazole (200 mg twice daily) treatment during 18 months.

One year after later, the patient presented again cough, mucopurulent sputum and weight loss. A chest X-ray showed greater thickening of the walls of the apical cavities. Small oval yeast-like cells were again disclosed, but in sputum smears stained by Grocott; and again *H. capsulatum* was isolated from sputum cultures. The patient was then treated with itraconazole (100 mg/day) for 6 months; in spite of his recovery the drug was maintained for more 7 months.

Follow-up: on October 95 the chest X-ray showed only fibrotic scars (Fig. 6) and the patient remained in good conditions 3 years after stopping antifungal therapy (November 96).

**COMMENTS**

Eleven cases of Chronic Pulmonary Histoplasmosis (CPH) were reported in Brazil (Table 1). The rarity of the diagnosis of CPH in Brazil can be explained partly by its clinical and radiological similarities with tuberculosis. On the other hand, mycological diagnostic facilities are not available in many places of the vast territory of Brazil. Therefore, pointing out that CPH is overlooked.

The first step to recognize CPH is to consider it in the differential diagnosis. Serologic test can be used as screening test for histoplasmosis; immunodiffusion is the best because complement fixation crossreaction does occur. However, *H* band sometimes may not be present. Mycological diagnosis may be based on the observation of the fungus in clinical specimens and its isolation and identification in culture.°

Expectoration is an usual symptom in CPH, so sputum is the easiest specimen to be examined. However, *H. capsulatum* may be recognized if sputum is smeared and stained by Gomori-Grocott technique. Nevertheless, fungus cells may be scarce or another yeast-like elements (*Candida* spp.) may be present, leading to misdiagnosing it (Fig. 2). For that reason, culture, using proper media and proper incubation temperature for the isolation and recognition of the fungus, is necessary.
TABLE 1
Eleven cases of chronic pulmonary histoplasmosis reported in Brazil.

<table>
<thead>
<tr>
<th>Case # (Ref)</th>
<th>Patient age, sex</th>
<th>Radiologic features</th>
<th>Material</th>
<th>Diagnosis M</th>
<th>C</th>
<th>I</th>
<th>IDh</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 (7)</td>
<td>41, M</td>
<td>Mass lesion in middle lobe</td>
<td>Lung (open lung biopsy)</td>
<td>+</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>2 (1)</td>
<td>45, M</td>
<td>“Delicate shadows at the inner aspects and bases of the lungs”</td>
<td>Sputum</td>
<td>+</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>3 (1)</td>
<td>35, M</td>
<td>“Discrete patch” at the apex of the right lung</td>
<td>Sputum</td>
<td></td>
<td>+</td>
<td></td>
<td></td>
</tr>
<tr>
<td>4 (9)</td>
<td>72, M</td>
<td>Bilateral lung consolidation and pleural effusion</td>
<td></td>
<td>H, M</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>5 (9)</td>
<td>53, F</td>
<td>Nodular lesions in both lungs and cavity in right lung</td>
<td></td>
<td>M</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>6 (2)</td>
<td>64, M</td>
<td>Reticulonodular infiltration in both lungs and cavities in the right upper lobe</td>
<td>Sputum</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>H, M</td>
</tr>
<tr>
<td>7 (2)</td>
<td>58, M</td>
<td>Reticulonodular infiltration in right lung and cavities in left lung</td>
<td>Sputum</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>H, M</td>
</tr>
<tr>
<td>8 (2)</td>
<td>74, M</td>
<td>Reticulonodular infiltration in both lungs, consolidation and pleural thickness in right upper lobe</td>
<td>Sputum</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>H, M</td>
</tr>
<tr>
<td>9 (3)</td>
<td>34, F</td>
<td></td>
<td>Sputum</td>
<td>–</td>
<td>+</td>
<td></td>
<td>H, M</td>
</tr>
<tr>
<td>10 (3)</td>
<td>42, F</td>
<td></td>
<td>Sputum</td>
<td>–</td>
<td>+</td>
<td></td>
<td>H, M</td>
</tr>
<tr>
<td>11 (3)</td>
<td>64, M</td>
<td></td>
<td>Sputum</td>
<td>–</td>
<td>+</td>
<td></td>
<td>H, M</td>
</tr>
</tbody>
</table>

Diagnosis: M = microscopy, C = culture, I = animal inoculation. IDh: immunodiffusion test to histoplasmosis.

Clinical specimen of CPH can be obtained with transthoracic needle biopsy. Tissue sections (case 1) or necrotic material (case 2) of part of the specimen may be used for microscopic examination, another part for culturing the fungus.

Two and half years after been treated with antifungals, patient (case 1) presented a fungus ball by A. fumigatus. The diagnosis was suggested by chest X-ray, presumed by immunodiffusion and proved mycologically. Active lesions of histoplasmosis were also detected in the patients lesions. This is the first case reported in Brazil, presenting such association.

RESUMO
Histoplasmosse pulmonar crônica no Brasil: relato de dois casos com cavitación diagnosticados por biópsia transtorácica

Apresentam-se dois casos de Histoplasmosse Pulmonar Crônica e outros onze casos da literatura brasileira são comentados. Após cura clínica, um de nossos pacientes apresentou bola fúngica asperrilg intracavitária, na parede da cavidade foi identificado H. capsulatum. Comentam-se aspectos diagnósticos da micose.
ACKNOWLEDGEMENTS

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