Pleural Effusion: Aetiology, Clinical Presentation and Mortality Outcome in a Tertiary Health Institution in Eastern Nigeria – A Five Year Retrospective Study

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Abstract

Aim: This retrospective notes review determined the aetiology, clinical presentation and mortality in patients with pleural effusion over a 5-year period.

Method: A retrospective audit of patients’ folders from January 2008-December 2012. Data collected included demographics, clinical presentation, and laboratory and mortality outcome.

Results: Of 199 folders reviewed, 108 were males. Male:female ratio was 1.18:1. Major symptoms were cough 156(78.4%), chest pain 142(71.4%) and dyspnoea 130(65.3%). Major signs included pyrexia 120(60.3%), ascites 48(24.1%) and hypotension 42(21.1%). The more common aetiology were TB 84(42.2%), parapneumonic 28(14.07%) and cardiac failure 28(14.07%). Forty-six (37.1%) of 120 patients screened were HIV positive. Mortality was 33(16.6%).

Conclusion: Pleural effusion is a common presentation in our clinical practice. Bacterial infection particularly TB is the most common cause. The mortality rate in patients with pleural effusion is still high. Determining the aetiology and early intervention are needed to reduce the mortality in patients with pleural effusion.

Keywords: Pleural effusion; Aetiology; Clinical features; Mortality; Eastern Nigeria

Introduction

Pleural effusion is a frequent clinical condition observed in pulmonary pathology. It also occurs in many other disease entities such as cardiac failure, liver diseases, renal diseases, endocrine disorders, malignancies and connective tissue diseases [1,2]. The clinical presentation varies from mild asymptomatic effusion to life threatening effusions with dyspnoea and chest pain.

Pleural effusion may be detected clinically when it is moderate to massive but only radiologically when it is mild. Chest x-ray is the commonest radiological approach used in detecting pleural effusion but ultrasound and computerized tomography scan of the chest are important in making a diagnosis. Qureshi et al. [3] in 2009 demonstrated the importance of thoracic ultrasound in differentiating malignant from benign pleural disease in patients presenting with suspected malignant pleural effusion. While Evans et al. [4] working in South Africa have demonstrated a high diagnostic yield of ultrasound guided trans-thoracic fine needle aspiration and ultrasound-assisted cutting needle biopsy in patients with superior vena cava syndrome with associated intrathoracic mass lesion. Cytology and laboratory evaluation of the pleural fluid is required to confirm the aetiologic cause of the effusion [1,5,6].

Aetiologic factors in pleural effusion have widely been studied in many African regions and beyond. The most common cause of the condition in low or moderate income countries has been attributed to infective causes and tuberculosis (TB) has remained the main cause in many areas; while in high income countries malignancy is a more common cause [7-10]. Its prevalence seems visible in West Africa (1.7-23%) [2,7]. In Western Nigeria, Onadoko et al. [11] found lymphocytic effusion in 76% of TB patients and 22% in patients with malignancy. Ezemba et al. [12] in Eastern Nigeria found non-specific pleurisy/empyema as the commonest cause of pleural effusion in a percutaneous needle biopsy of 37 patients. Sutherland et al. [6] in Gambia noted that TB was the commonest cause of pleural effusion but notoriously difficult to diagnose due to paucibacillary nature of the fluid. In Ghana the most common cause is TB with 63.5% of all effusions attributed to tuberculosis [10]. Desalew et al. [13] in Ethiopia noted that TB was the commonest cause of pleural effusion in that region with mortality in 6.4% of the patients. The work done in Cotonou Benin republic revealed 13% mortality in patients with pleural effusion. Likewise, a prospective study in Qatar revealed TB as the commonest cause of pleural effusion in that region with 32.5% of all causes, most patients being Asian immigrant workers [14]. Parasitic infections associated with pleural effusions have been documented; microfilaria have been detected in pleural effusions associated with malignancies [15-17].

Objective of the Study

Our study aimed at reviewing the aetiology and clinical presentation of patients with pleural effusion and the mortality outcome of these patients in a five-year retrospective study in a tertiary health institution in the South Eastern Nigeria.

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Methodology

This was a retrospective study which reviewed all in-patients seen with pleural effusion from January 2008 to December 2012. All in-patients folders seen during the period were selected. The folders were cross-checked with the register in the radiology Department of the Federal Medical Centre Owerri. Uncompleted folders were removed from the data. A form for data collection was created. The data collected included demographics (age, sex, location of effusion and year in review); clinical (medical history, symptoms, and physical signs elicited); laboratory (haemogram, microbiology, biochemistry, serology, cytology, radiology) and mortality outcome. Pleural effusion was radiologically classified as right hemithorax, left hemithorax and bilateral using Chest X-ray. They were also classified as exudates (pleural fluid protein>30 g/dL) or transudates (pleural fluid protein<30 g/dL).

Ethical Clearance

The Ethical clearance was obtained from The Ethics Committee of Federal Medical Centre Owerri, Imo State Nigeria.

Data Analysis

Data was analyzed using SPSS version 16 Chicago IL.

Results

During the period, 214 patients’ folders were reviewed. Fifteen folders were excluded due to incomplete data. One hundred and ninety-nine folders were analyzed. The mean age was 51 years. The maximum age was 96 years while the minimum was 15 years. There were 108 males and 91 females with male to female ratio of 1.18:1 (Table 1).

Average age=51years, Range=81 years (15-96 years).

The major symptoms found in these patients were cough 156(78.4%), chest pain 142(71.4%), and dyspnoea 130(65.3%). Major signs elicited were pyrexia 120(60.3%), and ascites 48(24.1%). Hypotension 42(21.1%), trachea Deviation 36(18.1%) and hypothermia 12(6.0%) (Table 2).

Medical history revealed tobacco and alcohol use in 66(33.2%) and 54(27.1%) respectively. 32(16.1%) had been treated for TB while 10(5.02%) had been treated for TB while 14(7.1%) of the 120 patients screened for HIV were positive. (Table 2).

The annual case presentation of pleural effusion increased steadily from 24 in 2008 to 56 in 2012. The commonest causes were infective with 46(37.1%) of the 120 patients screened. (Table 2).

Overall mortality recorded was 16.6%. Mortality remained at 29.2 in 2008 to 14.7% in 2009; then with slight reduction in 2012 (Table 3). Malignancy was the commonest cause of death. The primary location of malignancy was the breast in 28%, the liver in 12%, the lung in 10% and lymphoma in 7%.

The macroscopic appearance of thoracentesis fluid was documented in 118 patients. It was haemorrhagic in 49%, straw coloured in 31% and purulent in 20%. Pleural effusion was exudative in 82%.

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Table 3: Year specific mortality rate, Aetiology and mortality in patients with pleural effusion

<table>
<thead>
<tr>
<th>Year</th>
<th>Frequency (%)</th>
<th>Year specific mortality rate No (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>2008</td>
<td>24 (12.1)</td>
<td>7 (29.2)</td>
</tr>
<tr>
<td>2009</td>
<td>34 (17.1)</td>
<td>5 (14.7)</td>
</tr>
<tr>
<td>2010</td>
<td>40 (20.1)</td>
<td>6 (15.0)</td>
</tr>
<tr>
<td>2011</td>
<td>45 (22.6)</td>
<td>7 (15.6)</td>
</tr>
<tr>
<td>2012</td>
<td>56 (28.1)</td>
<td>8 (14.3)</td>
</tr>
<tr>
<td>Total</td>
<td>199 (100.00)</td>
<td>33 (100.00)</td>
</tr>
</tbody>
</table>

Discussion

Pleural effusion is a common presentation in medical and surgical practice both in low and middle income countries and high income countries. The cause of pleural effusion is primarily infective in low and middle income nations and malignant in the high income nations [2,7,8]. Most studies showed a tendency of more presentation in males [2,14]. Our study supported this with a slight male preponderance. The reason for this was attributed to more males who indulge in drugs

![Table 1: Age and Sex distribution.](image-url)
Mortality in patients with pleural effusion was high 33(16.6%). This is higher than that recorded in Ethiopia (6.4%) [13] and in Benin Republic (13%) [2]. The year specific mortality rate was highest in 2008 with 7(29.2%) recorded for that year. This remained fairly stable over a period of three years with slight drop in 2012 to 8(14.3%) table 3. This may not be readily explained but is thought to be due to early diagnosis and improvement in level of care. The highest mortality was found among our patients with malignancy, heart failure, TB/ HIV co-infection and chronic kidney diseases. The reason is not far- fetched since most patients with malignancies present late and lack of equipped oncology centre in our sub-region. This calls for a need for a well equipped oncology centre with surgeons knowledgeable in the management of malignant pleural effusion with full application of the BTS guidelines for the management of malignant pleural effusion [8]. However, it is also worthy of note that malignant pleural effusion is associated with high mortality rate even in developed countries since it is a sign of metastatic spread [24,25]. The patients with heart failure were mostly elderly people and probably with other co-morbidities. The increase in mortality in TB patients was high in most of the patients who were on retreatment for TB and patients with TB/HIV co-infection due to the immune compromised state and other co-morbidities. It was noted that these patients had other complications of tuberculosis such as chronic obstructive pulmonary disease and lung collapse. There was also marginal anaemia in most of the patients which further compounded the problem. There is therefore need for early detection of TB, early initiation of anti-tuberculous chemotherapy and close monitoring of TB chemotherapy using directly observed treatment (DOT). This will in turn reduce complications of TB including pleural effusion and also reduce mortality.

Implication for Future Research

The findings in this study shows that in our environment pleural effusion has variable aetiologic factors with tuberculosis and parapneumonic effusion topping the list. Doctors practicing in our sub region should have these differentials in mind as they are treatable causes. Further investigations such as pleural biopsy and histology should be considered to diagnose other conditions like malignancy to ensure early and appropriate therapy. The use of polymerase chain reaction, like the Gene-Xpert machine for diagnosis of mycobacterium tuberculosis and rifampicin resistance should be readily employed to enhance case detection of TB. The study is a retrospective study. A prospective study will be required to confirm some of the findings in this research work.

Conclusion

Pleural effusion is a common clinical presentation in our clinical practice with variable aetioloogy: Bacterial infections particularly tuberculosis remain the commonest aetiology. The mortality rate in patients with this condition is still high. Early investigation to determine the aetiology and appropriate early intervention is needed to reduce the mortality in patients with pleural effusion.

Acknowledgement

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References


