ROLE OF PLEURAL FLUID CHOLESTEROL IN DIFFERENTIATING TRANSUDATIVE AND EXUDATIVE PLEURAL EFFUSION

Tariq Mahmood1, Kaleem Ahmad2, Arvind Kumar Verma3, A. D. Shukla4, Abhinav C5, Srinivasa6

1Professor and Head, Department of Pulmonary Medicine, MLN Medical College, Allahabad, U. P.
23rd Year Junior Resident, Department of Pulmonary Medicine, MLN Medical College, Allahabad, U. P.
33rd Year Junior Resident, Department of Pulmonary Medicine, MLN Medical College, Allahabad, U. P.
4Associate Professor, Department of Pulmonary Medicine, MLN Medical College, Allahabad, U. P.
5Senior Resident, Department of Pulmonary Medicine, SGPGI, Lucknow.
6Senior Resident, Department of Pulmonary Medicine, KMC, Mangalore.

ABSTRACT

BACKGROUND

Pleural effusion is defined as an abnormal and excessive collection of fluid in the pleural space. The most common cause in the West and India are infections followed by malignancy. In India, tubercular effusion is the most common cause followed by malignant effusion and a very few due to parapneumonic effusion.

Aims and Objectives- To study the role of pleural fluid cholesterol in differentiating transudative and exudative pleural effusion and comparison of pleural fluid cholesterol to Light’s criteria.

MATERIALS AND METHODS

In this prospective study, 204 patients with pleural effusion were included. Pleural fluid cholesterol, total protein, lactate dehydrogenase (LDH) as well as serum total protein and LDH levels along with other required investigations were studied. Clinical classification of transudative or exudative was done on the basis of aetiology.

RESULTS

Based on pleural fluid examination for cholesterol, protein, LDH and other haematological investigations and response to treatment besides history, clinical examination finding and chest radiograph, 173 patients were classified as exudates and 31 as transudates. Using pleural fluid cholesterol levels at a cut-off point of greater than 45 mg/dL for distinguishing transudates and exudates the sensitivity, specificity, positive predictive value (PPV) and negative predictive value (NPV) were 98%, 100%, 98% and 100% respectively. In the present study when pleural fluid cholesterol cut-off was taken as 60 mg/dL it was found that the sensitivity of differentiating exudates was 82%, while specificity was found to be 100% with PPV of 100% and NPV of 44%. Using Light’s criteria for discriminating transudates and exudates, sensitivity, specificity, PPV and NPV were found to be 98%, 100%, 100% and 92% respectively. In the present study, pleural fluid protein > 3 gm/dL when used as a differentiating parameter for transudates and exudates had a sensitivity of 98%, specificity and PPV of 100% and NPV of 74% respectively.

CONCLUSION

Pleural fluid cholesterol is cost-effective and useful parameters in distinguishing transudative from exudative aetiologies with the advantage of requiring only one laboratory determinations and no simultaneous blood sample compared to the use of Light’s criteria.

KEYWORDS

Pleural Fluid Cholesterol, Transudative, Exudative, Light’s Criteria.


BACKGROUND

Pleural effusion is defined as an abnormal collection of fluid in the pleural space. Two types of effusions can develop (transudative and exudative). Transudative pleural effusions are caused by fluid leaking into the pleural space and increased hydrostatic pressure or decreased osmotic pressure. Exudative effusions are caused by blocked blood vessels, inflammation, lung injury and drug reactions which causes damage or disruption of pleural membranes or vasculature (Barter et al, 1994).

The most common cause in the West are infections followed by malignancy,1 while in India it is tubercular effusion followed by malignant effusion2 and a very few due to parapneumonic effusion.

Exudative pleural effusions are a common diagnostic problem in clinical practice as the list of causes is quite exhaustive,1 although sometimes they can be inferred from the clinical picture. The aetiological distribution of pleural effusions in various series depends on the geographical area, patient’s age and advances in the diagnostic methods and treatment of the underlying causes. The difficulty in determining the cause of pleural effusion is shown by the fact that in many series “unknown aetiology” constitutes nearly 15%.3 Exudative effusions require to be separated into infectious causes, non-infectious causes and malignancy.
Aims and Objectives
- To study the diagnostic value of pleural fluid cholesterol in differentiating transudative and exudative pleural effusion.
- To compare pleural fluid cholesterol level for exudates with Light’s criteria.
- To evaluate the usefulness of diverse combinations of pleural cholesterol concentration, pleural or serum protein and lactate dehydrogenase (LDH) levels for the differentiation of pleural exudates and transudates.

Materials and Methods
This study was conducted in the Department of Pulmonary Medicine, MLN Medical College, Allahabad, over a period of 1 year from June 2016 to July 2017. All patients of either sex with pleural effusions were enrolled in this study as per inclusion and exclusion criteria. This was an observational cross-sectional study.

A total of 204 patients (153 males and 51 females) with clinical diagnosis of pleural effusion were enrolled for study. Among 204 patients 173 patients were diagnosed as exudative pleural effusion (tubercular, para-pneumonic and malignancy), while 31 patients were cases of Transudative pleural effusion (Congestive heart failure, Chronic liver disease etc.).

All patients qualifying for inclusion criteria were explained in detail about the study and procedures involved and informed consent was taken. A detailed history was taken about symptoms such as cough, expectoration, breathlessness, fever, weight loss, haemoptysis and chest pain. Clinical examination and necessary investigations were done including chest radiograph postero-anterior view, sputum acid-fast bacilli stain, USG thorax, pleural fluid aspiration, and analysis for protein, sugar, LDH, cholesterol, ADA, cell count and required blood investigations. The statistical analysis was performed using Microsoft Excel. The sensitivities, specificities, positive predictive values and negative predictive values were obtained. The aetiological classification according to the criteria of Light et al was used as the “Gold Standard.”

Results
A total of 204 patients (153 males and 51 females) with clinical diagnosis of pleural effusion were enrolled for study. Among 204 patients 173 patients were diagnosed as exudative pleural effusion (tubercular, para-pneumonic and malignancy), while 31 patients were cases of transudative pleural effusion (Congestive heart failure, Chronic liver disease etc.).

Study was carried out in a time duration of June 2016 to July 2017. The commonest type of effusion was found to be Tuberculous (158 cases) followed by Congestive heart failure (21 cases), Malignancy (11 cases), Para-pneumonic (4 cases), Chronic liver disease (5 cases), Rheumatic heart disease (4 cases) and one case of Nephrotic syndrome was identified as shown in Table 1. The most frequent cause of pleural exudates is tuberculosis followed by lung cancer, which is similar to the result of a study done in high tuberculosis burden countries.

<table>
<thead>
<tr>
<th>Diagnosis</th>
<th>Transudate</th>
<th>Exudate</th>
</tr>
</thead>
<tbody>
<tr>
<td>Tuberculosis</td>
<td>5</td>
<td>153</td>
</tr>
<tr>
<td>Congestive heart failure</td>
<td>13</td>
<td>8</td>
</tr>
<tr>
<td>Malignancy</td>
<td>1</td>
<td>10</td>
</tr>
<tr>
<td>Chronic liver disease</td>
<td>5</td>
<td></td>
</tr>
<tr>
<td>Para-pneumonic effusion</td>
<td></td>
<td>5</td>
</tr>
<tr>
<td>Nephrotic syndrome</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>Rheumatic heart disease</td>
<td>4</td>
<td></td>
</tr>
</tbody>
</table>

Table 1. Clinical Diagnosis of Patients Present with Pleural Effusion according to Light’s Criteria

Percentage Distribution of Disease
176 (84%) were exudates and 28 (16%) cases were transudates according to Light’s criteria. (Pleural fluid protein/serum protein ratio greater than 0.5, pleural fluid LDH/serum LDH ratio greater than 0.6, pleural fluid LDH level greater than two-thirds the upper limit of the laboratory’s reference range of serum LDH). It was found that the sensitivity and specificity of separating transudate and exudates with Light’s criteria is 98% and 100% respectively.

Percentage Distribution of Exudates and Transudates
Out of total 204 patients, 173 had pleural fluid cholesterol level greater than 45 mg/dL with sensitivity 98% and specificity 100%, and 135 patients had cholesterol level greater than 60 mg/dL with sensitivity of 82% and specificity of 100%. When pleural fluid cholesterol alone was used at a cut-off point of 45 mg/dL, 3 out of the 176 (as compared to Light’s criteria) was misclassified as transudate (sensitivity 98%), while there was no misclassification of 28 transudates (specificity 100%).
When pleural fluid cholesterol with a cut-off point of greater than 60 mg/dL was used as criteria for exudates, 31 out of 176 patients were misclassified as transudative pleural effusion, (sensitivity 82%) and 135 patients had exudative pleural effusion, while there is no misclassification of transudative pleural effusion with specificity of 100%.

The mean value of pleural fluid cholesterol in tuberculous, malignant and transudative pleural effusion was found to be 82.7, 66.6 and 27.3 respectively.

Comparison of Cholesterol Cut-Off Criteria

When pleural fluid protein and serum protein ratio > 0.5 alone was used as a criterion, 156 out of 204 patients classified had exudative pleural effusion with sensitivity of 81% and specificity of 100%. Pleural fluid LDH and serum LDH ratio > 0.6 alone classified 177 exudate out of 204 patients with specificity of 88% and sensitivity of 100%.

### Table 2. Shows Sensitivity, Specificity, Positive Predictive Value (PPV) and Negative Predictive Value (NPV) of various Laboratory Parameters for Transudates and Exudates Separation

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Sensitivity %</th>
<th>Specificity %</th>
<th>PPV %</th>
<th>NPV %</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pleural fluid cholesterol cut-off &gt; 45 mg/dL</td>
<td>98</td>
<td>100</td>
<td>98</td>
<td>100</td>
</tr>
<tr>
<td>Pleural fluid cholesterol cut-off &gt; 60 mg/dL</td>
<td>82</td>
<td>100</td>
<td>100</td>
<td>44</td>
</tr>
<tr>
<td>Pleural fluid protein/serum protein &gt; 0.5</td>
<td>81</td>
<td>100</td>
<td>100</td>
<td>44</td>
</tr>
<tr>
<td>Pleural fluid LDH Serum LDH ratio &gt; 0.6</td>
<td>100</td>
<td>88</td>
<td>97</td>
<td>100</td>
</tr>
<tr>
<td>Light's criteria</td>
<td>98</td>
<td>100</td>
<td>100</td>
<td>91</td>
</tr>
<tr>
<td>Pleural fluid LDH &gt; 2/3 UL of Serum LDH</td>
<td>82</td>
<td>100</td>
<td>100</td>
<td>46</td>
</tr>
<tr>
<td>Pleural fluid protein &gt; 3 gm/dL</td>
<td>94</td>
<td>100</td>
<td>100</td>
<td>74</td>
</tr>
</tbody>
</table>

In this study when pleural fluid protein ≥ 3 gm/dL was used to separating transudates and exudates 158 patients had exudative pleural effusion, while 46 patients were identified as having transudative pleural effusion with sensitivity 94%, specificity 100%, positive predictive value 100% and negative predictive value of 74%.

When pleural fluid LDH > 2/3 of upper limit of serum LDH used as parameters it has 82% sensitivity, 100% specificity positive predictive value of 100% and negative predictive value of 46%.

On Pearson correlation test pleural fluid cholesterol correlation is 1 and on the basis of protein ratio this is 0.9. This suggests that pleural fluid cholesterol is highly correlated than protein ratio with clinical diagnosis for exudates for level of significance at the level of α = 0.01.

For level of significance (α = 0.05), for pleural fluid cholesterol > 45 mg/dL and pleural fluid cholesterol > 60 mg/dL, p = 0.08 which is greater than level of significance (α = 0.05). This shows that there is no significance between these two cut-off criteria.

### Table 3. Mean Values of ADA in Tuberculous and Non-Tuberculous Pleural Effusion

<table>
<thead>
<tr>
<th>Studies</th>
<th>Tuberculous</th>
<th>Non-Tuberculous</th>
</tr>
</thead>
<tbody>
<tr>
<td>SK Sharma et al⁵</td>
<td>95.8</td>
<td>30.7</td>
</tr>
<tr>
<td>PK Sinha et al⁶</td>
<td>76.8</td>
<td>14.5</td>
</tr>
<tr>
<td>Dharmendra A Bhamaniya et al⁷</td>
<td>65.48</td>
<td>24.29</td>
</tr>
<tr>
<td>Present Study</td>
<td>62.33</td>
<td>24.33</td>
</tr>
</tbody>
</table>

### Table 3. Shows Sensitivity, Specificity, Positive Predictive Value (PPV) and Negative Predictive Value (NPV) of various Laboratory Parameters for Transudates and Exudates Separation

DISCUSSION

Pleural effusion is commonly encountered condition in clinical practice. The initial step in the management of pleural effusions is to distinguish transudates from exudates. Various biochemical tests have been used in diagnosis of pleural effusion. This study has evaluated the diagnostic value of different biochemical tests used for evaluation of pleural fluid.

The criteria often used to do so are based on biochemical parameters proposed by Light et al.⁴ Since no single test has yet proved to be completely satisfactory, the search for improved methods continues.

In the present study, majority of pleural effusions were of exudative nature (86%). As the most of cardiac, renal and
liver disease patients first approach the Internal Medicine Department of hospital, most of the transudative effusions were managed further by Medicine Department, so the number of transudative effusions managed in Pulmonary Medicine Department of the hospital were relatively less. In this study, tubercular pleural effusion was found in 78% of patients.

Table 2 shows comparison of different studies for sensitivity, specificity, PPV and NPV of various biochemical parameters.

The cholesterol levels are elevated in exudative pleural effusions of much shorter duration. The cause of the increased cholesterol concentration in pleural exudates is unknown. Increased pleural permeability leading to accumulation of cholesterol in pleural exudates due to “serum leakage” may be a reasonable explanation. Cholesterol is found in all tissues and is uniformly found in all pleural effusions.

Our results show that an increased concentration of cholesterol greater than 45 mg/dL in pleural fluid constitute useful measurements for separating exudates from transudates. This finding of our study is also supported by previous study of Heffner et al.

In our study we found that in transudative, parapneumonic, tubercular and neoplastic pleural effusions, pleural fluid cholesterol levels with a classifying threshold of 45 mg/dL has a sensitivity of 98 percent and specificity of 100 percent for diagnosis of exudates with a PPV of 100.

In the present study when pleural fluid cholesterol cut-off was taken as 60 mg/dL, it was found that the sensitivity of differentiating exudates was 82%, while specificity was found to be 100% with PPV of 100% and NPV of 44%. Pleural fluid to serum cholesterol ratio had a sensitivity of 88% and a specificity of 100% with PPV of 100% and NPV of 56%.

Dharmendra A Bamaniya et al. in his study found that 86.2% of patients with TB and 79.16% of malignancy patients had pleural fluid cholesterol levels > 60 mg/dL with pleural fluid to serum cholesterol ratio > 0.4.

Guleria R et al. investigated the role of pleural fluid cholesterol in differentiating transudative from exudative pleural effusion. Pleural fluid cholesterol had a sensitivity of 88% and a specificity of 100% for exudates with an accuracy of 92% with cut-off value of 60 mg/dL or above.

Anand K et al. in his study using pleural fluid cholesterol levels at a cut-off point of greater than 60 mg/dL and/or total protein at a cut-off point of greater than 3 mg/dL for distinguishing transudates and exudates reported the sensitivity, specificity, positive predictive value (PPV) and negative predictive value (NPV) were 100 percent.

AB Hamal et al. reported that pleural fluid to serum protein ratio greater than 0.5 has a sensitivity of 81.4% and specificity of 82.6%; pleural fluid to serum LDH ratio has a sensitivity of 86% and specificity of 94.7% and pleural fluid cholesterol with sensitivity of 97.7% and specificity of 100% for differentiating exudative and transudative pleural effusions.

In our study pleural fluid cholesterol criterion correctly classified eight cases of CHF as Transudative effusion, which were misclassified as exudative effusion by Light’s criterion. It was most probably due to diuretic therapy frequently given in CHF treatment. In our study it was found that mean value of pleural fluid ADA was 62.33 in tubercular effusion and 24.33 in non-tubercular effusion which is comparable to other studies as shown in Table 3, hence high levels of ADA is more diagnostic towards tuberculosis. Pleural fluid ADA is a specific and a sensitive marker to differentiate between tuberculous and non-tuberculous pleural effusion and it is becoming popular and a vital tool to differentiate tuberculous from non-tuberculous cases of pleural effusion which is quite handful in countries like India where tuberculosis is prevalent everywhere.

In present study pleural fluid protein > 3 g/dL when used as a differentiating parameter for transudates and exudates had a sensitivity of 98%, specificity and PPV of 100% and NPV of 74% respectively. Present study is also supported by Anand K et al. which shows sensitivity of 98%, specificity of 100%, PPV of 100% and NPV of 92%.

In our study pleural fluid cholesterol and serum cholesterol ratio has sensitivity, specificity, PPV and NPV of 88%, 100%, 100% and 44% respectively, which was comparable to Lights et al. and John Heffner et al.

Pleural fluid LDH level > 2/3 of upper limit of serum LDH reported sensitivity, specificity, PPV and NPV of 82%, 100%, 100% and 46% respectively, which was comparable to Lights et al. and John Heffner et al study.

Our study also shows that the most common cause of exudative pleural effusion in India is tuberculosis.

CONCLUSION

The present study entitled “Role of pleural fluid cholesterol in differentiating transudative and exudative pleural effusion” included 204 patients.

The Conclusions of this Study were as follows:

- Pleural fluid cholesterol is simple, cost effective and useful parameter for differentiation of transudates from exudates with a cut-off level of 45 mg/dL.
- Measurement of pleural fluid total protein also has a good sensitivity and specificity in differentiation of exudative and transudative effusion.
- Our study supports other studies in establishing the vital role and cost effectiveness of these novel biochemical markers (pleural fluid total protein and pleural fluid cholesterol) in comparison to long list of markers in evaluation of pleural effusion, and hence recommends their use.
- Our criteria of using pleural fluid cholesterol to differentiate between exudative and transudative pleural effusion is found to be at par with Light’s criteria, which continues to be the most sensitive and specific parameter.
- Pleural fluid cholesterol can be used to correctly classify the type of effusion in case of congestive heart failure with diuretic therapy where light’s criteria may be misleading.
- The most common cause of exudative pleural effusion in India is tubercular.
- The mean value of pleural fluid cholesterol in tuberculous, malignant and transudative pleural effusion was found to be 82.7, 66.6 and 27.3 respectively.
In present study, mean value of ADA level was 62.33 in tubercular effusion and 24.23 in non-tubercular effusion which is comparable to other studies as shown in Table 3. This suggests that high levels of ADA in pleural fluid is more diagnostic towards tuberculosis.

REFERENCES


