

Study of Etiology of Pleural Effusion in Andhra Pradesh PopulationYasar Arafath Shaik¹, Dadeboyina suryakala²¹Assistant Professor, Department of General Medicine, Kurnool Medical College, Kurnool, Andhra Pradesh-518002.²Assistant Professor, Department of General Medicine, Kurnool Medical College, Kurnool, Andhra Pradesh-518002.

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Conflict of interest: Nil

Abstract**Background:** Pleural effusion is an excessive accumulation of fluid in the pleural space. It can be a diagnostic dilemma for clinicians because of the multiple etiologies of lung, pleura or systemic disorders.**Material and Methods:** 95 patients with newly diagnosed pleural effusion based on Chest – x-ray PA view were studied. 20 ml of pleural fluid was aspirated and sent for biochemical, microbiological, and pathological analysis. Echocardiography, USG abdomen, and biopsy of the pleura were carried out in the same patients whose etiology or diagnosis was not clear.**Results:** Clinical manifestations were: 65 (68.4%) had fever, 75 (78.9%) had cough, 44 (46.3%) had breathlessness, 22 (23.1%) had pedal oedema, 46 (48.4%) had chest pain, and 5 (5.26%) had abdominal distention. 57 (60%) patients had tubercular pleural effusion, and 38 (40%) had non-tubercular pleural effusion. In non-tubercular pleural effusion 9 had (23.6%) synpneumonic effusion, 6 (15.7%) had CCF, and 13 (34.2%) had malignancy, 2 (2.1%) had rheumatoid arthritis, 2 (2.1%) had dengue fever, 2 (2.1%) had pancreatitis, and 4 (4.20%) had nephrotic syndrome.**Conclusion:** In the present study, it was concluded that most of the pleural effusions are tubercular, but the etiological evaluation of pleural effusions is very important to diagnose and treat accordingly.**Key words:** tubercular, non-tubercular, malignant, cell cytology, CCF, cirrhosisThis is an Open Access article that uses a funding model which does not charge readers or their institutions for access and distributed under the terms of the Creative Commons Attribution License (<http://creativecommons.org/licenses/by/4.0>) and the Budapest Open Access Initiative (<http://www.budapestopenaccessinitiative.org/read>), which permit unrestricted use, distribution, and reproduction in any medium, provided original work is properly credited.**Introduction**

Pleural effusion is an excessive accumulation of fluid in the pleural space, it indicates an imbalance between pleural fluid formation and removal [1]. Accumulation of pleural fluid is not a specific disease, but rather a reflection of the underlying pathology [2]. Therefore, for a patient with pleural effusion, it is a challenge for clinicians to know the exact etiology of pleural effusion. Without knowing the cause of pleural effusion, it can't be treated appropriately. Pleural effusions accompany a wide variety of disorders of the lung, pleura and systemic disorders [3].

Hence to diagnose pleural effusion appropriately, pleural fluid cytology, biochemistry and clinical presentation are important to establish an etiological diagnosis. The common causes of pleural effusion are CCF, cirrhosis, nephrotic syndrome, tuberculosis, malignancy. Hence attempt is made to evaluate the etiology of pleural effusion and treated accordingly.

Material and Methods

95 adult patients who visited the medicine department of Kurnool Medical College Hospital in Kurnool, Andhra Pradesh-518002 were studied.

Inclusion criteria: Adult patients aged between 18 to 65 years in whom pleural effusion was newly diagnosed.

Exclusion criteria: patients who were critically ill, age <18 and >65 years were excluded from the study

Methods : Chest X-ray PA view was taken. All the patients were subjected to diagnostic thoracentesis. Under aseptic precautions, about 20 ml of fluid was aspirated and subjected to pleural fluid analysis. Biochemical, microbiological, and pathological analyses were done. Apart from this AFB stain and sputum for AFB, routine blood examination and ESR were also studied. Echocardiography, USG abdomen done based on clinical manifestations.

Duration of study: May 2023 to October 2023

Statistical Analysis: The patients having similar clinical manifestations were classified by percentage. Tubercular and non-tubercular patients were segregated by percentage. Non-tubercular patients were noted with different etiologies and classified by percentage. The ratio of males and females was 2:1.

Observation and Results

Table 1: Clinical manifestations of patients with pleural effusion 65 (68.4%) had fever, 75 (78.9%)

had cough, 44 (46.3%) had breathlessness, 22 (23.1%) had pedal oedema, 46 (48.4%) had chest pain, and 5 (5.26%) had abdominal distention.

Table 1: (Total No of patients- 95) Clinical manifestations of patients with pleural effusion

Sl no	Clinical manifestations	No of patients	Percentage %
1	Fever	65	68.4
2	Cough	75	78.9
3	Breathlessness	44	46.3
4	Pedal edema	22	23.1
5	Chest pain	46	48.4
6	Abdominal distention	5	5.26

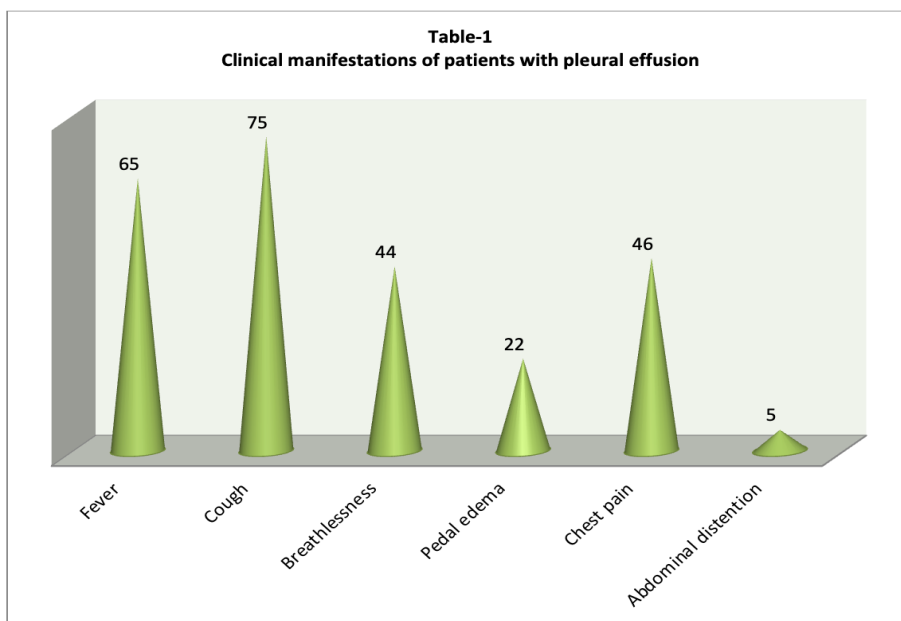


Table 2: Types of plural effusion 57 (60%) had tubercular plural effusion, and 38 (40%) had non-tubercular plural effusion.

Table 2: Types of pleural effusion

Sl no	Particulars	No of patients	Percentage %
1	Tubercular pleural effusion	57	60
2	Non- tubercular pleural effusion	38	40

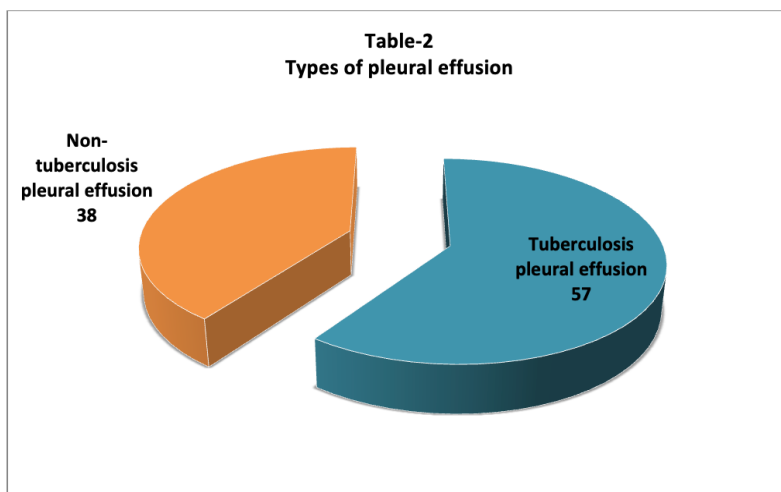
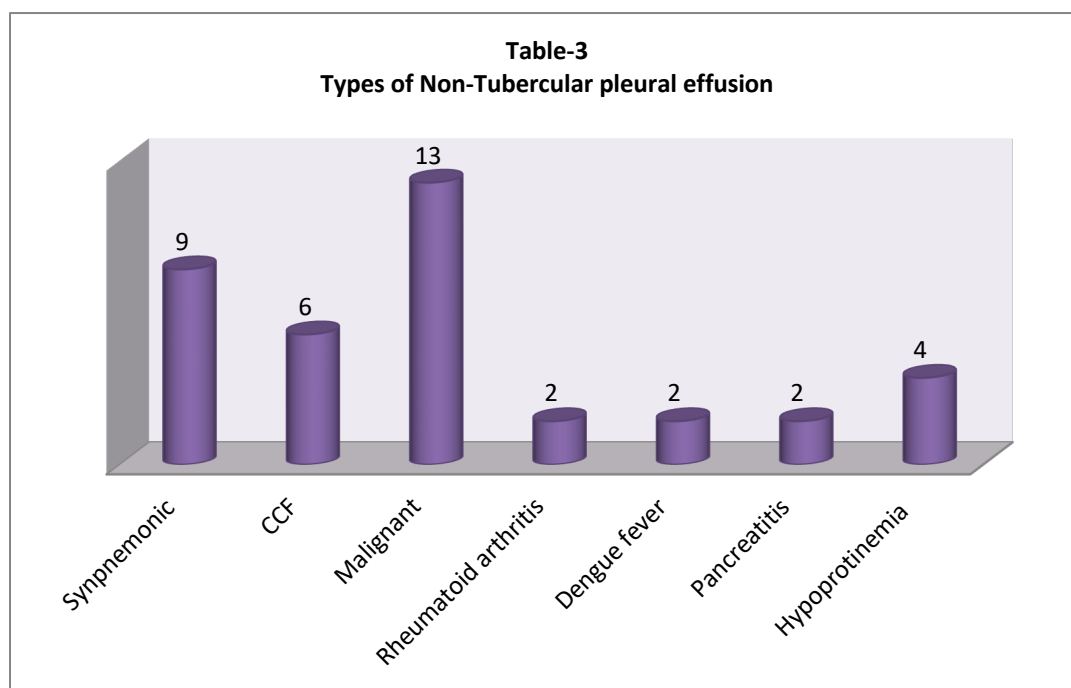


Table 3: Types of non-tubercular pleural effusion (out of 38 non-tubercular effusions): 9 (23.6%) synpneumonic effusion, 6 (15.7%) CCF, 13 (34.2%) had malignant effusion, 2 (2.1%) had rheumatoid arthritis, 2 (2.1%) had dengue fever, 2 (2.1%) had pancreatitis ,4 (4.21%) had nephrotic syndrome.

Table 3: Types of Non-Tubercular pleural effusion (No of patients 38)

Sl no	Particulars	No of patients (38)	Percentage %
1	Synpneumonic	9	23.6
2	CCF	6	15.7
3	Malignant	13	34.2
4	Rheumatoid arthritis	2	2.1
5	Dengue fever	2	2.1
6	Pancreatitis	2	2.1
7	Nephrotic syndrome	4	4.21



Discussion

Present study of pleural effusion in the Andhra Pradesh population 65 (68.4%) had a fever, 75 (78.9%) had a cough, 44 (46.3%) had breathlessness, 22 (23%) had pedal oedema, 46 (48.4%) had chest pain, and 5 (5.26%) had abdominal distention (Table 1). Out of 95 patients, 57 (60%) had tubercular pleural effusion and 38 (40%) were non-tubercular (Table 2). Among the non-tubercular, 9 (23.6%) were synpneumonic effusion, 6 (15.7%) had CCF, 13 (34.2%) had malignancy, 2 (2.1%) had rheumatoid arthritis, 2 (2.1%) had dengue fever, 2 (2.1%) had pancreatitis, and 4 (4.21%) had nephrotic syndrome (Table 3). These findings were more or less in agreement with previous studies [4-7].

The history of the patients provides information about the possible aetiology of pleural effusion (PE) and guidelines for necessary investigations. A history of pneumonia suggests para-pneumonic effusion, either complicated (empyema-like) or uncomplicated; fever indicates an infective etiology. A

history of cardiac, renal, or liver impairment can suggest transudative effusion; older age, weight loss, and a history of smoking point towards a diagnosis of malignant Pleural Effusion. Recent pedal edema, or deep venous thrombosis (DVT), may result in an effusion related to pulmonary embolism. Trauma may result in hemothorax or chylothorax. Recent esophageal procedures or a history of alcohol use suggest Pleural effusion related to oesophageal rupture. Physical findings such as ascites may indicate cirrhosis, ovarian cancer, or Meigs syndrome. 75% of malignant pleural effusions (PE) are caused by neoplasms of the lung, breast, ovary, lymphoma or mesothelioma. [7] Inflammatory pleural effusion is an uncommon complication seen in about 2% to 5% of patients with rheumatoid arthritis. [8] Pancreatitis related pleural effusion is largely due to the close proximity of the pancreas to the diaphragm. [9] patients with CCF and pleural effusion present with orthopnea and paroxysmal nocturnal dyspnea and on lung auscultation have crackles. Pleural effusion is associated with cirrhosis of the liver and is called

Hepatic hydrothorax. Effusion is caused by the passage of ascetic fluid from the peritoneal cavity into the pleural space through diaphragmatic defects [10-12]. About 20% of patients with nephrotic syndrome develop pleural effusion from severe hypoalbuminemia, which leads to decreased oncotic pressure.

Tuberculosis is quite common in underdeveloped countries due to poor socio-economic conditions.

Summary and Conclusion

The present study of the aetiology of pleural effusion in the Andhra Pradesh population will be quite helpful to physicians and radiologists as pleural effusion (PE) is a common clinical problem. The major cause of pleural effusion may be tuberculosis, followed by pneumonic and malignant causes, cirrhosis of the liver, etc. The most common cause of pleural transudate was CCF. Histological examination and pleural biopsy were most useful to find out the exact cause of Pleural effusion in both tubercular and malignant effusions. But this study demands further pathophysiological, nutritional, genetic, and immunological studies because the exact factors and pathogenesis of pleural effusion are still unclear.

Limitations of Study: Owing to the tertiary location of the research centre, the small number of patients, and the lack of the latest techniques, we have limited findings and results.

- This research paper was approved by the ethical committee of Kurnool Medical College Hospital, Kurnool, Andhra Pradesh (51802).
- No conflict of interest
- Self funding

References

1. Kays MD Pleuropulmonary complications of pancreatitis. *Thorax* 1968, 23, 297–305.
2. Albert WM, Salem AJ- Hepatic hydrothorax cause and management *Arch Intern Med* 1991, 151, 2383–88
3. A Yat sen H, - Diaphragmatic defect as a cause of massive hydrothorax in cirrhosis of the liver, *Med.* 1998, 18, 216–20
4. Cavina C, Vicini G – Radiological aspects of pleural effusion in medical nephropathy in children, *Ann Radiol Diagn (Bologna)*, 1958, 31, 163-202. Italian.
5. Chinchkar NJ, Talwar. D, Jain SK. – A surprise approach to the etiologic diagnosis of pleural effusion in the respiratory intensive care unit and short-term evaluation of treatment lung India, 2015, 32, 95–112.
6. Mehta AA, Patel MN, - investigation into the Role of Medical Pleuroscopy in the Diagnosis and Management of Patients with Pleural Diseases, *Ind. J Thoracic. Carivasc. Surg.* 2012, 28,120–6.
7. B Ijazk, Mohd. KT- management of tuberculosis pleural effusion. *J Biomed SC. and Res.* 2011.3.302
8. Y Khan, M. Alsamwi. M- Etiology of Pleural Effusion among Adults in the State of utar: A 1-Year Hyear Study *Eastern Mediterranean Health Journal* 2011, 17(7), 611-618
9. Light R.W- clinical practice, pleural effusion *New England Journal of Medicine* 2002, 346, 1971–77
10. Marel M – The incidence of pleural effusion in a well-defined region *Epidemiologic study in central biochemia, Chest* 1993, 104, 1486–89
11. Mudaly DK, Deo SVS, subi TS- an update in the management of malignant pleural effusion *Ind. J. Palliat Care* 2011, 17, 98–101
12. Walker WC, Wright V- pulmonary lesions and rheumatoid arthritis *medicine*, 1968, 47, 501–520.