

Non-malignant Pleural Effusions

Umair Gauhar, MBBS
Clinical Instructor
Division of Pulmonary, Allergy, Critical Care
and Sleep Medicine
Ohio State University Medical Center

Outline

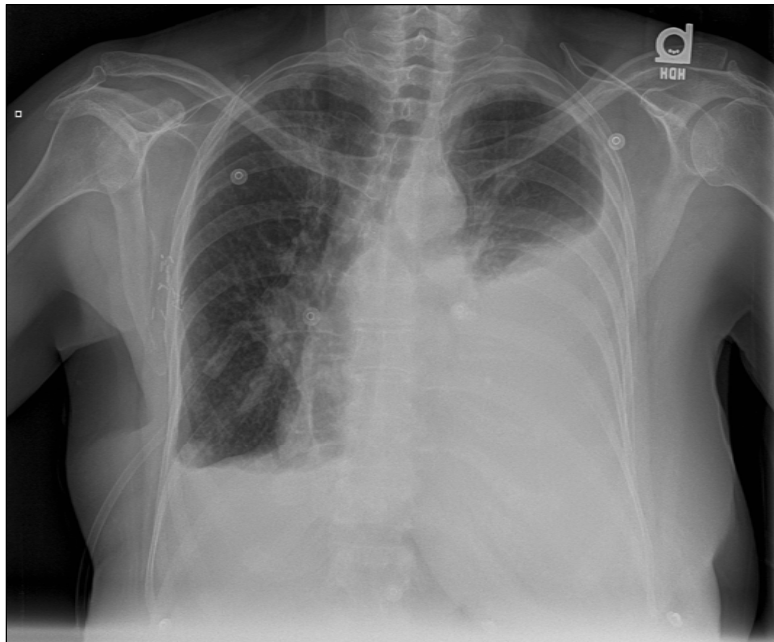
- **Case presentation**
- **Epidemiology**
- **Pathophysiology**
- **Diagnostic approach**
 - **Pleural fluid analysis**
- **Parapneumonic effusions/ empyema**
- **Treatment**

Case presentation

- 65 years old woman presents with 3 days of
 - Fever
 - Malaise
 - Cough
 - Purulent sputum
 - Worsening dyspnea
 - Left sided pleuritic chest pain
- Past Medical/Surgical History
 - COPD
 - Hypertension
 - Congestive Heart Failure
 - Cholecystectomy
- Social history
 - Current smoker
 - 50 pack-years

Case presentation (contd.)

- Physical examination
 - Febrile 101.5°F
 - Tachypneic
 - Tachycardiac
 - Hypoxemic (O2 sats 88% on RA)
 - Decreased breath sounds and tactile fremitus on left
- Lab data
 - WBC 22,000
 - Neutrophils 90%
 - Sodium 129
 - Chest x-ray (shown)



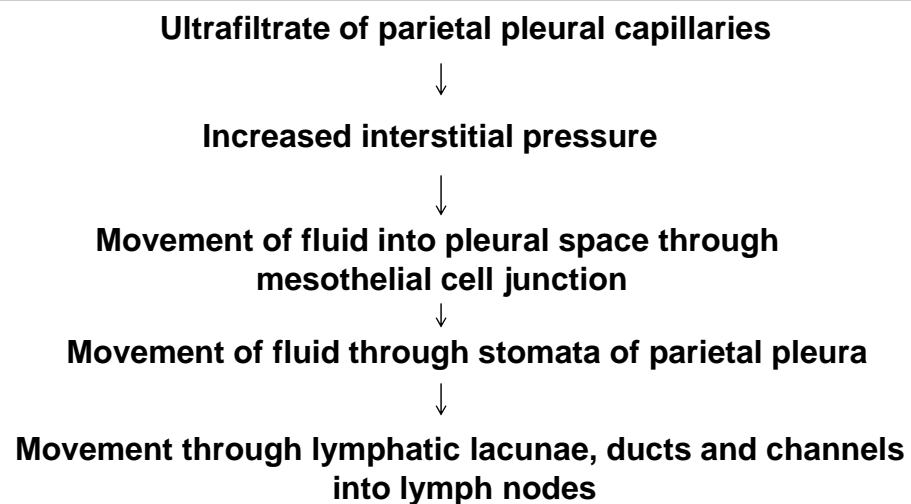
Epidemiology

- **Estimated 1.5 million cases of pleural effusions in the United States annually**
 - Light R. *Pleural Diseases*. 4th ed. Philadelphia, PA: Williams and Wilkins; 2001
- **Associated with a wide variety of diseases**
- **Congestive heart failure, pneumonia and malignancy accounting for two thirds of the cases**

Pathophysiology

- **Parietal pleura supplied by microvessels from intercostal artery**
 - **Located close to mesothelial surface**
- **Visceral pleura supplied by microvessels from bronchial circulation**
 - **Located at a distance from the mesothelial surface**
- **Normal pleural fluid formation/ resorption are functions of the parietal pleura**

Pathophysiology (contd.)



Pathophysiology (contd.)

- **Normal pleural fluid**
 - **0.1 to 0.2 ml/kg**
 - **Clear**
 - **Low protein (1.0 to 1.5 g/dl)**
 - **< 1500 nucleated cells / μ L**
 - **61% to 77% monocytes-macrophages**
 - **9 to 30% mesothelial cells**
 - **7% to 11% lymphocytes**
 - **2% neutrophils**
 - **0% eosinophils**
 - **pH \geq 7.60**

Pathophysiology (contd.)

- **Mechanism of abnormal pleural fluid formation**
 - **Increased hydrostatic pressure (CHF)**
 - **Decreased oncotic pressure (hypoalbuminemia)**
 - **Decreased pleural pressure (trapped lung)**
 - **Increased endothelial permeability (pneumonia)**
 - **Decreased lymphatic drainage (malignancy)**
 - **Movement from peritoneal space (hepatic hydrothorax)**
 - **Movement from extra-vascular space (duropleural fistula, migrated/ misplaced CVC/ feeding tube)**

Diagnostic Approach

1. Clinical History

- Could be asymptomatic
- Dyspnea and chest pain are the two most common presenting symptoms
- Dyspnea most likely from
 - decreased chest wall compliance
 - depression of ipsilateral diaphragm and
 - increased output from neurogenic receptors
- Dyspnea out of proportion to exam findings can suggest PE

Diagnostic Approach

1. Clinical History

- Chest pain
 - Usually pleuritic
 - Intensity proportional to degree of pleural inflammation
 - May be decreased by splinting by manual pressure over the chest wall
 - May be localized or radiating
 - Central diaphragmatic inflammation causes radiating pain in the posterior neck, shoulder and trapezius area

Diagnostic Approach

1. Clinical History

Asymptomatic

- BAPE
- Rheumatoid pleural effusion
- Nephrotic syndrome
- Yellow nail syndrome
- Trapped lung
- Urinothorax
- Peritoneal dialysis associated effusion

Symptomatic

- Bacterial pneumonia
- Lupus pleuritis
- Postcardiac injury syndrome
- Pulmonary embolism
- Congestive heart failure

Diagnostic Approach

1. Clinical History

▪ Useful clues

- Orthopnea, PND, lower extremity → CHF
- H/O asbestos exposure → BAPE
- H/O alcoholism, poor dentition, loss of consciousness → aspiration/ anerobic empyema
- H/O of CABG → post-cardiac injury syndrome or trapped lung
- H/O retching → esophageal rupture
- H/O SLE (or procainamide use) → lupus pleuritis
- Obstructive uropathy → urinothorax
- Spinal surgery or trauma → duropleural fistula

Diagnostic Approach

2. Physical Examination

- Signs depend on volume of pleural effusion
 - < 300 ml → not detectable on physical examination
 - 500 ml → dull percussion, decreased fremitus, decreased breath sounds
 - > 1000 ml → bulging of ICS, decreased chest expansion, bronchovesicular sounds and egophony at upper level of effusion

Diagnostic Approach

3. Chest Radiograph

- Sensitivity proportional to volume of pleural fluid
 - 5ml → blunting of posterior costophrenic angle on lateral decubitus film
 - 50-75 ml → blunting of posterior costophrenic angle on lateral view
 - 175-200 ml → blunting of costophrenic angle on PA film
 - > 500 ml → opacification of lung base



Diagnostic Approach

3. Chest Radiograph

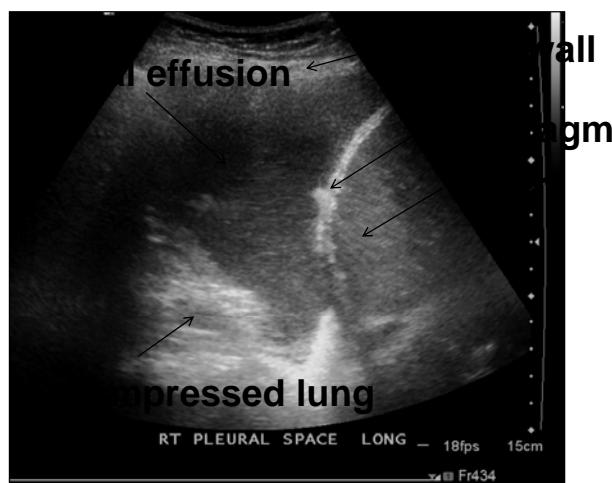
- **Non-malignant causes of massive effusion with mediastinal shift**
 - Tuberculosis
 - Empyema
 - Hepatic hydrothorax
 - Chylothorax
 - Hemothorax
 - Congestive heart failure

Diagnostic Approach

4. Ultrasound

1. Diagnosis and sampling of loculated pleural effusions

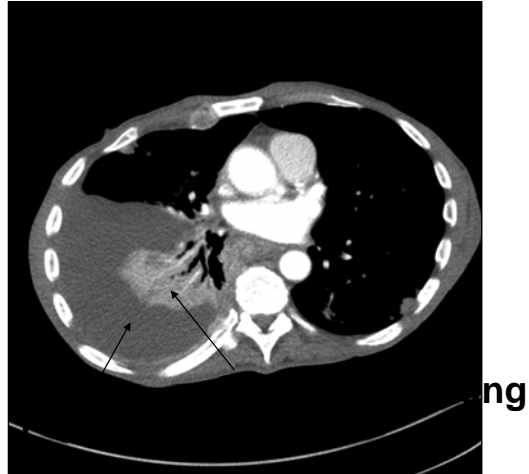
2. Guided sampling of small pleural effusions



Diagnostic Approach

5. Computed Tomography

- Most sensitive radiographic study
- Useful for differentiating free flowing effusions, loculated effusions, parenchymal lesions and extrapleural disease



Diagnostic Approach

6. Pleural Fluid Analysis (PFA)

- Observation acceptable in
 - Small effusions (< 1 cm thickness on lateral decubitus films)
 - Patients presenting with typical symptoms of CHF and bilateral pleural effusions of similar size and absence of chest pain or fever
- PFA should be performed in all new effusions
- Therapeutic thoracentesis (1 to 1.5 L) in symptomatic effusions

Diagnostic Approach

6. Pleural Fluid Analysis (PFA)

- Prospective study of 129 patients (Chest 1987; 91:817-822)
 - PFA provided definitive diagnosis in 18% of patients
 - Presumptive diagnosis in 55% of patients
 - Non-diagnostic in 27% of patients
 - Non-diagnostic PFA helpful in excluding infections
- Approximately 30 ml of fluid needed for a complete PFA

Diagnostic Approach

6. Pleural Fluid Analysis (PFA)

- Diagnoses that can be established by PFA
 - Empyema → pus
 - TB pleuritis → + AFB smear and/ or culture
 - Fungal disease → + KOH stain or culture
 - Lupus pleuritis → high PF ANA or LE cells
 - Chylothorax → chylomicrons
 - Hemothorax → PF/ blood ratio > 0.5
 - Biliopleural fistula → PF bilirubin/ serum > 1.0

Diagnostic Approach

6. Pleural Fluid Analysis (PFA)

- **Diagnoses that can be established by PFA (contd.)**
 - Peritoneal dialysis associated effusion → protein < 0.5 g/dL, PF/serum glucose > 1.0
 - Esophageal rupture → high salivary amylase, low pH, food particles in PF
 - Rheumatoid pleurisy → low glucose (< 30 g/dL), low pH, debris
 - Extravascular migration of CVC or feeding tube
 - Urinothorax → PF/ serum Cr > 1.0
 - Duropleural fistula → $\beta 2$ transferrin in pleural fluid

Diagnostic Approach

6. Pleural Fluid Analysis (PFA)

- **Observation of pleural effusion**
 - Pale yellow (straw colored) → transudate, some exudates
 - Red (bloody) → BAPE, PCIS, pulmonary infarction, trauma
 - White (milky) → chylothorax, cholesterol effusion (satin like sheen)
 - Brown → chronic bloody, ruptured amebic liver abscess (anchovy paste)
 - Black → *Aspergillus niger*
 - Yellow-green → rheumatoid pleurisy (with debris)
 - Color of enteral feeds or IV infusate → Feeding tube or CVC has entered pleural space
 - Pus → empyema
 - Putrid → anerobic empyema
 - Urine → urinothorax
 - Water like → duropleural fistula

Diagnostic Approach

6. Pleural Fluid Analysis (PFA)

- **Transudate vs. Exudate**
 - **Exudate if anyone of the following criteria met**
 - **PF protein > 2.9 g/dL**
 - **PF/S protein > 0.5**
 - **PF LDH > 0.67 upper limits serum LDH**
 - **PF/S LDH > 0.6**
 - **PF cholesterol > 45 mg/dL**
 - **PF/S cholesterol > 0.3**
 - **Serum albumin-PF albumin < 1.2 g/dL**

Am J Respir Crit Care Med. 1995; 151: 1700-1708

Diagnostic Approach

6. Pleural Fluid Analysis (PFA)

- **Transudates**
 - **CHF (most common cause)**
 - **Hepatic hydrothorax**
 - **Atelectasis**
 - **Nephrotic syndrome**
 - **Hypoalbuminemia**
 - **Trapped lung**
 - **Peritoneal dialysis associated pleural effusion**
 - **Urinothorax (only transudate with a pH < 7.30)**
 - **Duropleural fistula (β 2-transferrin present in PF)**

Diagnostic Approach

6. Pleural Fluid Analysis (PFA)

- **Exudates**
 - Infections (bacterial, viral, fungal, parasitic, mycobacterial)
 - Pulmonary embolism (can be transudative)
 - Post CABG
 - Post cardiac injury syndrome (PCIS)
 - GI disease associated (pancreatitis, esophageal rupture)
 - Connective tissue diseases (SLE, RA, WG, CSS, SS)
 - BAPE

Diagnostic Approach

6. Pleural Fluid Analysis (PFA)

- **Exudates (continued)**
 - Sarcoidosis
 - Uremia
 - Meig's syndrome
 - Endocrinopathies (hypothyroidism, ovarian hyperstimulation syndrome)
 - Yellow nail syndrome
 - Drug reaction (amiodarone, nitrofurantoin, dantrolene, methotrexate)
 - Lung entrapment
 - Radiation therapy
 - Chylothorax
 - Hemothorax

Diagnostic Approach

6. Pleural Fluid Analysis (PFA)

- **Pleural fluid glucose and pH**
 - Direct relationship
 - Increased metabolism by neutrophils and bacteria → increased CO₂, lactic acid (empyema, esophageal rupture)
 - Poor efflux of CO₂ and lactate due to pleural membrane thickening (rheumatoid pleural effusion)
- **Low PF glucose and pH**
 - Complicated parapneumonic effusion/ empyema
 - Esophageal rupture
 - Tuberculous empyema
 - Chronic rheumatoid pleurisy
 - Lupus pleuritis

Diagnostic Approach

6. Pleural Fluid Analysis (PFA)

- **Cell count and differential count**
 - Total cell count nonspecific
 - Differential count more helpful
 - Neutrophil predominant (> 50% of nucleated cells)
 - Parapneumonic (with infiltrate)
 - PE (without infiltrate)
 - Acute viral infection
 - Pancreatitis
 - Acute TB pleuritis (rarely)

Diagnostic Approach

6. Pleural Fluid Analysis (PFA)

- Cell count and differential count
 - Lymphocyte predominance (> 80% of nucleated cells)
 - TB pleurisy
 - Chylothorax
 - Yellow nail syndrome
 - Chronic rheumatoid effusion
 - Sarcoidosis
 - Acute lung rejection
 - Uremic pleural effusion
 - Post-CABG surgery

Diagnostic Approach

6. Pleural Fluid Analysis (PFA)

- Cell count and differential count
 - Eosinophilic predominant (> 10% of nucleated cells)
 - Pneumothorax
 - Hemothorax
 - BAPE
 - Pulmonary infarction
 - Parasitic disease
 - Fungal disease
 - Drug induced lung disease
 - *Unlikely to be TB pleuritis*

Diagnostic Approach

6. Pleural Fluid Analysis (PFA)

- **PF amylase**
 - Acute pancreatitis (pancreatic)
 - Pancreatic pseudocyst -several thousand fold (pancreatic)
 - Esophageal rupture (salivary)
 - Pneumonia (salivary)
 - Ruptured ectopic pregnancy (salivary)

Diagnostic Approach

6. Pleural Fluid Analysis (PFA)

Chylothorax

- Acute/ subacute
- Trauma, surgery, LAM
- Dyspnea common
- Milky/ turbid or bloody
- Lymphocyte predominant exudate
- PF Tg >110 mg/dL, PF Tg < 50 mg/dL excludes chylothorax
- PF cholesterol < 200mg/dL
- Chylomicrons in PF

Pseudochylothorax

- Chronic, insidious
- RA, TB, empyema
- Dyspnea uncommon
- Milky, satin like sheen
- Neutrophil predominant exudate
- PF Cholesterol > 200mg/dL
- PF Tg maybe > 110mg/dL
- Cholesterol crystals in PF

Diagnostic Approach

6. Pleural Fluid Analysis (PFA)

- PF rheumatoid factor and ANA
 - Not routinely performed
 - PF ANA > 1:320 or greater than serum ANA → lupus pleuritis
 - PF rheumatoid factor > 1:320 or greater than serum RF → rheumatoid pleurisy
 - Can be high in bacterial pneumonia
- PF adenosine deaminase (ADA)
 - 40-60 units/ L → TB pleurisy
 - PF/serum ADA > 1 → TB, RA, empyema

Diagnostic Approach

6. Pleural Fluid Analysis (PFA)

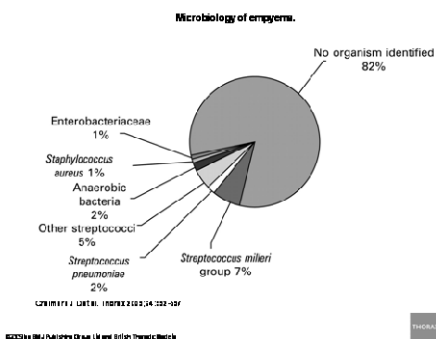
- Microbiological tests (cultures, stains)
 - Positive Gram stain or culture diagnostic of empyema
 - TB pleuritis
 - Pleural biopsy histology 63% to 85% sensitive
 - Pleural biopsy culture 55% to 80%
 - Pleural fluid culture 13% to 70%
 - Pleural biopsy smear 5% to 18%
 - Pleural fluid AFB smear < 5%

Parapneumonic Effusions/ Empyema

- 40% to 57% of cases of pneumonia associated with parapneumonic effusion
- Complicated parapneumonic effusions in 10% to 15% of patients
- Empyema (pus in the pleural space) in 5% of patients
- Exudative phase (0–72h)→ fibrinopurulent phase (3–10 days)→ organizational phase (10–21 days)

Parapneumonic Effusions/ Empyema (contd.)

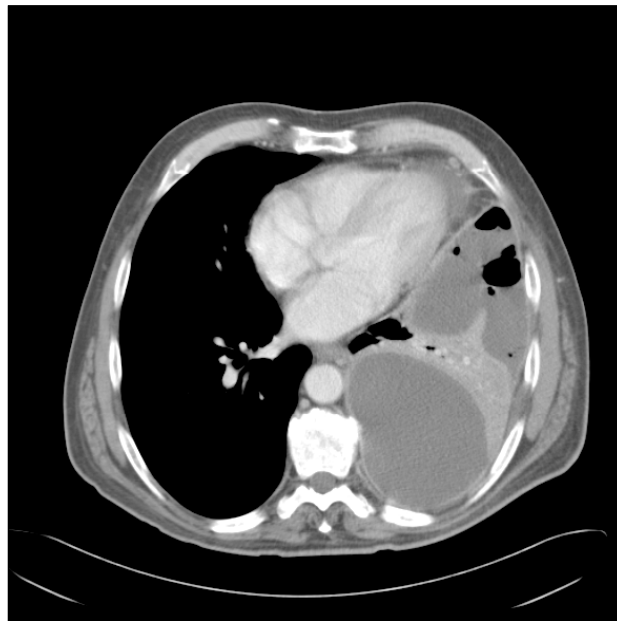
- Microbiology
 - Anerobic bacteria
 - *S. pneumoniae*
 - *Staphylococcus aureus*
 - *H. influenza*
 - *Klebsiella pneumoniae*
 - Gram negative bacilli
 - Fungi (incidence increasing)
 - Atypical organisms, virus, parasites



Parapneumonic Effusions/ Empyema (contd.)

▪ Diagnosis

- Exudative to gross pus
- Cell count may be low due to cell lysis
- pH < 7.30
- Low glucose
- High LDH
- + Gram stain and/or culture
- CT chest with contrast showing “split pleura sign” with pleural enhancement
- Pleural fluid loculations (seen better with chest ultrasound)
 - Indicates poor prognosis



Parapneumonic Effusions/ Empyema (contd.)

Table 1—Categorizing Risk for Poor Outcome in Patients With PPE

Pleural Space Anatomy		Pleural Fluid Bacteriology		Pleural Fluid Chemistry*	Category	Risk of Poor Outcome	Drainage
A ₀ minimal, free-flowing effusion (< 10 mm on lateral decubitus)†	AND	B _X culture and Gram stain results unknown	AND	C _X pH unknown	1	Very low	No‡
A ₁ small to moderate free-flowing effusion (> 10 mm and < ½ hemithorax)	AND	B ₀ negative culture and Gram stain§	AND	C ₀ pH ≥ 7.20	2	Low	No
A ₂ large, free-flowing effusion (≥ ½ hemithorax)¶ loculated effusion, # or effusion with thickened parietal pleura**	OR	B ₁ positive culture or Gram stain	OR	C ₁ pH < 7.20	3	Moderate	Yes
		B ₂ pus			4	High	Yes

*pH is the preferred pleural fluid chemistry test,⁸ and pH must be determined using a blood gas analyzer.^{9,10} If a blood gas analyzer is not available, pleural fluid glucose⁸ should be used (P₀ glucose ≥ 60 mg/dL; P₁ glucose < 60 mg/dL). The panel cautions that the clinical utility and decision thresholds for pH and glucose have not been well-established.

†Clinical experience indicates that effusions of this size do not require thoracentesis for evaluation, but will resolve.²

‡If thoracentesis were performed in a patient with A₀ category pleural anatomy and P₁ or B₁ status found, clinical experience suggests that the P₁ or B₁ findings might be a false-positive. Repeat thoracentesis should be considered if effusion enlarges and/or clinical condition deteriorates.

§Regardless of prior use of antibiotics.

¶If clinical condition deteriorates, repeat thoracentesis and drainage should be considered.

#Larger effusions are more resistant to effective drainage, possibly because of the increased likelihood that large effusions will also be loculated.¹¹

#Pleural loculations suggest a worse prognosis.¹²

**Thickened parietal pleura on contrast-enhanced CT suggests presence of empyema.¹³⁻¹⁵

Chest 2000; 18: 1158-1171

Treatment of Non-malignant Effusions (contd.)

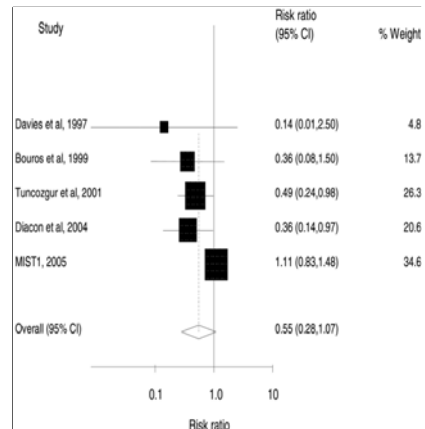
- Treat the underlying cause
- Persistent or recurrent effusions can require repeat thoracentesis, indwelling pleural catheter, tube thoracostomy or pleurodesis
- Hepatic hydrothorax→ medical management, TIPS
 - VATS to repair diaphragmatic defect
 - Chemical pleurodesis rarely successful
 - Chest tube contraindicated
- Lupus pleuritis→ steroids
- Trapped lung→ Reassurance if asymptomatic; decortication if symptomatic and underlying lung normal

Treatment of Non-malignant Effusions (contd.)

- Rheumatoid pleurisy → resolve spontaneously after several months
- PCIS → NSAIDs, steroids
- Chylothorax → hyperalimentation, medium chain TG, bed rest, thoracic duct ligation
- Parapneumonic effusions/ empyema → antibiotics, pleural drainage, intrapleural fibrinolytics (?), decortication (VATS vs. thoracotomy)

Treatment of Non-malignant Effusions (contd.)

- Intrapleural fibrinolytic therapy for empyema
- No evidence for routine use
- Trend towards reducing need for surgery in some groups
- Recommended by ACCP and BTS on a case to case basis



CHEST 2006; 129:783-790

Management of Malignant Pleural Effusion

Shaheen Islam, MD, MPH

Associate Professor

Director, Interventional Pulmonology

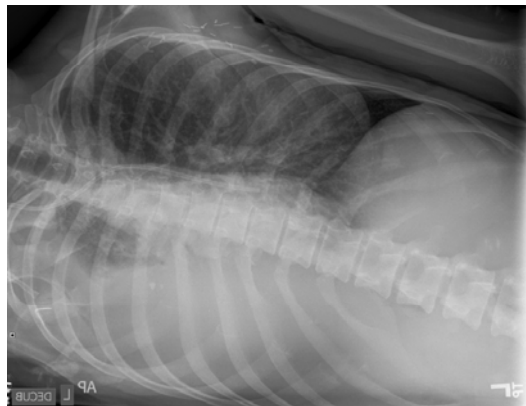
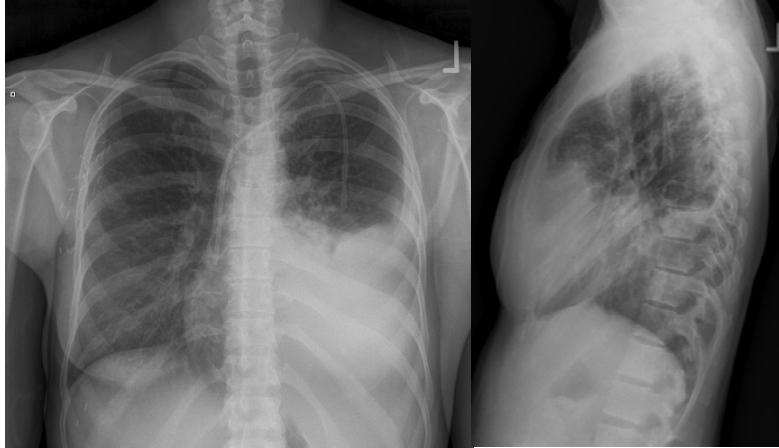
Assoc Medical Director, Pulmonary Diagnostics Lab

**Division of Pulmonary, Allergy, Critical Care
and Sleep Medicine**

Ohio State University Medical Center

Case

- **37 year old female with metastatic breast cancer and left pleural effusion**
- **1 week history of dyspnea**



Epidemiology of MPE

- **Most common exudative effusion 40%~70%**
- **Lung 32%, Breast 18%, Lymphoma 11%**
- **More common in females <60 (50% vs 35%)**
- **15% of patients with Lung Cancer have MPE at diagnosis**

Gomez et al. CHEST 2007, 618S
Marel M. Eur Respir Mon, 2002;22:146-156

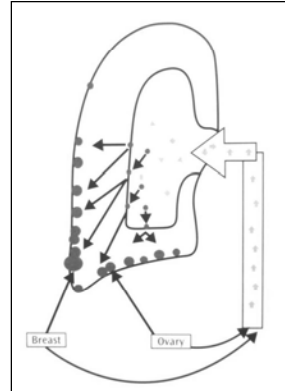
Epidemiology

- **Primary tumor not identified 5-10% MPE**
- **20-30% of malignant lymphoma**
- **50% of patient with breast Ca**
- **Ovarian, GI, mesothelioma ~15%**
- **Usually <6 months survival except in breast Ca**

Marel M. Eur Respir Mon, 2002;22:146-156
Ruckdeschel JC Semin Oncol 1995; 22:58-63

Etiology

- Direct invasion
- Increased capillary permeability
- Tumor emboli to visceral pleura
secondary seeding of parietal pleura
- Hematogenous spread to parietal pleura



Paramalignant effusion

- Effusion without pleural involvement
- Low oncotic pressure
- Blockage of mediastinal lymphatics
 - Lymphoma, Squamous Cell Ca
- Postobstructive pneumonia, atelectasis
- Trapped lung
- Post radiation
- Chemo related

Statement by ATS on MPE, AJRCCM 2000; 162:1987-2001

Clinical features

- **Progressive Dyspnea**
 - **Decreased chest wall compliance**
 - **Mediastinal shift**
 - **Increased shunt fraction from atelactatic lung**
- **Dull chest pain**
 - **Malignant mesothelioma**
- **Cough**

Diagnosis

History

Symptoms

Imaging

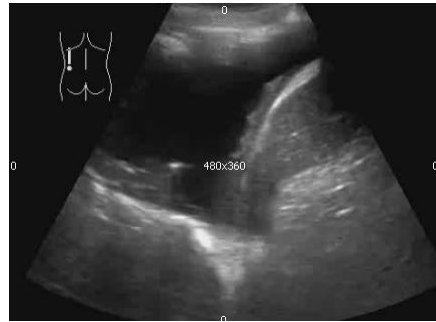
CXR, CT chest, Ultrasound

Thoracentesis

Pleuroscopy/Pleural biopsy

Case

- Therapeutic thoracentesis (1.8L removed)
- Bedside ultrasound by clinician
- Symptom relieved
- No PTX
- Biochemical studies ordered



Pleural Fluid results

- Protein 4.5/7.1
- LDH 645/219
- WBC 1653
 - Lymphocytes 15%
- Malignant cells 85%

Other Fluid Studies

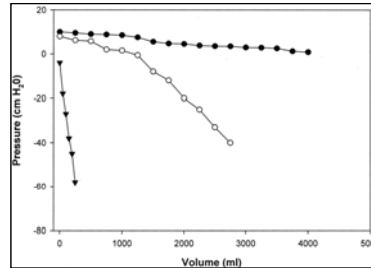
- **Amylase**
 - Lymphoma, Ovarian Ca, Pancreatic Ca
 - CEA, B72.3, Leu-M1
 - Calretinin and cytokeratin 5/6 identifies mesothelioma but not benign mesothelial cells
- **Flow cytometry**
 - if lymphocytic effusion with possibility of lymphoma
- **Tumor markers**
 - CEA (>10-12ng/mL)
 - Vascular Endothelial Growth Factor (VEGF)

Goal for thoracentesis

- **Diagnostic & therapeutic (large volume)**
- **Any relief of symptom?**
- **Will it recur?**
- **If so, how soon?**
- **Did the lung expand?**

Entrapped Lung?

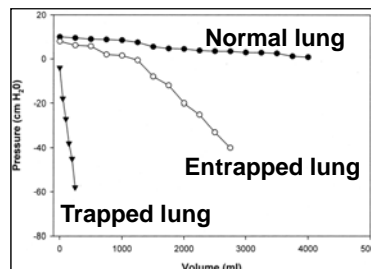
- Pleural Manometry
- Lung Elastance (Pel)
 - Change in pleural pressure in relation to volume of fluid removed <19cm H₂O/L
- Entrapped lung
 - Visceral pleural restriction from malignancy or active disease
 - poor success with pleurodesis
 - Pneumothorax ex-vacuo



Doelken et al. Chest 2004; 126:1764-69

Entrapped Lung?

- Pleural Manometry
- Lung Elastance (Pel)
 - Change in pleural pressure in relation to volume of fluid removed <19cm H₂O/L
- Entrapped lung
 - Visceral pleural restriction from malignancy or active disease
 - poor success with pleurodesis
 - Pneumothorax ex-vacuo



Doelken et al. Chest 2004; 126:1764-69

Patient Update



- Left lung expanded after thoracentesis
- Dyspnea improved
- However, effusion returned after 12 days

Management options

- Repeat thoracentesis
- Indwelling Pleural Catheter Placement
- Pleurodesis
 - Chest tube
 - Medical thoracoscopic
 - VATS
- Pleuroperitoneal shunts
- Surgery

What is the best option?

Personalized Care

Primary Tumor type

- **NSCLC effusion respond poorly to chemo**
- **Small cell Ca effusion respond to chemo**

Pleural Fluid Tests

- **Poor survival with**
 - **Low pH**
 - **Low glucose**
 - **High LDH**
 - **CEA**

Performance Status

- **Karnofsky score <30, <1 month survival**

Lung re-expansion

- **Extensive intrapleural deposition, multiple loculations, trapped lung, endobronchial airway obstruction will cause of failure of pleurodesis**
- **Pneumothorax after large volume thoracentesis suggest trapped lung**

Patient Preference

- **Duration of hospital stay**
- **Invasiveness of procedures**
- **Success of a definitive therapy**
- **Associated risks**

Individual Management Options

Repeat thoracentesis

- Only if reaccumulation >30 days
- Limited life expectancy
- Poor performance status
- May trigger cytokine, fibrin and cause loculations
- Large volume thoracentesis with pleural manometry safe

Chemical Pleurodesis

- **Dyspneic and life expectancy more than 4-6 weeks**
- **Frequent recurrence with symptoms**
- **Success rate 71%~97%**

Heffner J. Semin Respir Crit Care Med 2010; 31:723-733

Chemical Agents

- **Doxycycline**
 - **Severe CP, thru Chest tube**
 - **Used with lidocaine**
- **Talc:**
 - **Slurry thru CT**
 - **Lower success rate**
 - **Poudrage during thoracoscopy 90% success rate ***
 - **No ARDS with larger calibrated particles**
- **Other Agents**
 - **Quinacrine**
 - **Bleomycin**
 - **Silver nitrate**
 - **IFN alpha-2b**

Thoracoscopic vs Chest tube Pleurodesis

- No large RCT available
- Cochrane review of 112 patients
 - Slightly better with thoracoscopy
- Better success with thoracoscopy in breast and primary lung Ca
- Center dependent
- May be better with talc poudrage vs talc slurry

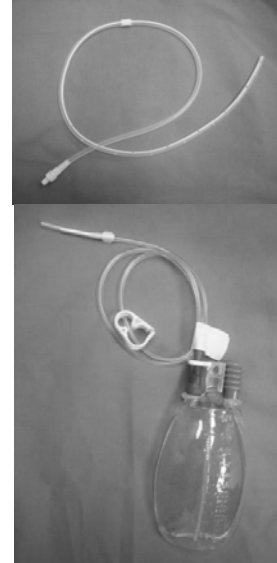
Shaw P et al. Cochrane database Systemic review 2004; CD002916
Dressler et al Chest 2005; 127:909-915

Medical Thoracoscopy vs. VATS

- Similar
- Adhesions if present can be lysed
- Poudrage sprayed effectively under vision
- Sedation
- VATS more invasive
- Cost
- Requires 3-7days of hospital stay

Indwelling Pleural Catheter

- Outpatient placement
- Patients may remain active with good QOL
- Can be placed in trapped lung or in failed pleurodesis cases
- Complications
 - Obstruction, tumor seeding, infection



Indwelling Pleural Catheter

- Outpatient placement
- Patients may remain active with good QOL
- Can be placed in trapped lung or in failed pleurodesis cases
- Complications
 - Obstruction, tumor seeding, infection



Indwelling Pleural Catheter

- **Spontaneous pleurodesis in 40%**
- **Removed in 60% with resolution of effusion**
- **Sclerosants can be instilled through catheter**

Putnam et al. Cancer 1999; 86:1992-1999
Musani et al. Respiration 2004;71:559-566
Warren et al. Eur J Cardiothoracic Surg 2008;33:89-94

Surgery

- **Pleuroperitoneal Shunting with 95% efficacy if other options fail**
- **Parietal pleurectomy**
- **Decortication**
- **Higher mortality**

Patient Update

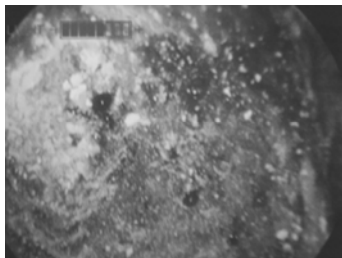


Examination of pleural space



Pleural space after talc poudrage

Thoracoscopic Pleurodesis



Conclusion

- **MPE indicates advanced disease**
- **Palliative management is variable depending on tumor type, patient preference, life expectancy, fluid characteristics, performance status and available resources**
- **Tunneled catheter is safe and cost effective**