Radiologically Guided Chest Tube Insertion with Fibrinolytic Instillation for Loculated Pleural Effusion/Empyema

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Abstract: Title; Radiologically guided chest tube insertion with fibrinolytic instillation for loculated pleural effusion/empyema

Background; The management of loculated pleural effusion and empyema by chest tube drainage usually fails because of thick viscous fluid and multiple pleural space loculations. The use of radiographic assisted chest tube drainage with intrapleural fibrinolytic agents facilitates pleural drainage and can obviate the need for more invasive surgical interventions in these types of effusions.

Objectives; to evaluate the role of radiographic chest tube drainage with intrapleural fibrinolytic therapy with streptokinase as an adjunctive therapy in the management of loculative pleural effusion and empyema

Material and methods; 40 patients of CPE and empyema were considered for radiographic assisted chest tube drain with adjunctive intrapleural fibrinolytic therapy. Intrapleural, STK was administered 12-24 hourly in the dosage of 2,500 IU in 100 ml of saline. The end points were volume of fluid drained and radiological resolution.

Results; Statistical analysis showed a success of 65% in study group and in thoracotomy group 97.5% with minimal complications in both the groups.

Conclusion; Patients of loculative pleural effusion/empyema should be first subjected to radiographic assisted chest tube drainage as we have seen from results and then add fibrinolytic if loculations present.

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I. Introduction:

Empyema thoracis is defined as pleural space suppurative fluid collection and is a disease entity that has been recognized throughout recorded medical history. The incidence of empyema has fallen since the introduction of antibiotics for the treatment of pulmonary infections. Early diagnosis and effective therapy are essential. The late complications of untreated empyema with a fixed fibrotic chest cavity should only remain in the history books. Pulmonary infections due to iatrogenic causes remain the most important etiology of the loculated pleural effusion/empyema because people now live so long with cardiac failure, rheumatic disorder and breast cancer1,2.

Dry pleurisy may be the first indication of pleural inflammation although this is quickly followed by the outpouring of fluid rich in protein and polymorphs. Although frank purulence should develop before an effusion is referred to as an empyema, a pH<7 is the best marker for the need for operative intervention, and therefore is probably the discriminator for empyema. Continued accumulation of pus compresses the lung with shift of mediastinum to the opposite side. Fibrin is continually deposited on pleural surfaces producing a thickened rind, the deeper layers of which become fibrotic and avascular. The established empyema is walled off and the space is fixed. This allows for open drainage or easy stripping of the empyema membrane (thickened rind) during surgical decortications. Such decortications at this stage will produce full re-expansion of the lung with gradual resolution of the pleural inflammation and no functional impairment. It is this fibrin deposition which limits the diffusion of oxygen that leads to anaerobic respirations and the fall in pH within the empyema.

If the pus is not drained effectively or the empyema cavity excised, the continual formation and fibrosis of the pleural rind progressively restricts chest wall and diaphragmatic movement, eventually producing a shrunken, flattened, immobile hemi-thorax with overlapping ribs and scoliosis to the affected side.

Sometimes empyema may push through the chest wall at one of the perforating pathways of the neurovascular bundle, forming a collar-stud abscess which may break down giving a discharging fistula in continuity with the empyema cavity (empyema necessitans)3.
II. Etiology:
Despite the availability of antibiotic, 50% of all empyema’s currently diagnosed are secondary to complications of a primary pneumonic process in the lungs. Other causes include spontaneous pneumothorax, chest trauma, sub phrenic abscess, retained foreign bodies in the bronchial tree, esophageal diseases or operations on the esophagus, pulmonary or breast malignancy and surgical procedures involving the lung and mediastinum.

Pulmonary infections which include-
Pneumonia-(most common)
Lung abscess
Fungal infections
Tuberculosis
Trauma
Chest injuries-including implanted foreign bodies
Post thoracotomy
Ruptured esophagus/leaking anastomosis
Trans-diaphragmatic spread;
Sub phrenic abscess
Hepatic amoebiasis
Iatrogenic; multiple aspirations for effusions
Osteomyelitis- ribs or vertebrae
Septicemia

Prior to development of effective antibiotic treatment, pneumococcus and streptococcus were the most frequent causative organism of empyema. Staphylococcus has now emerged as the most frequent causative organism, particularly in children <2years, in whom it is cultured in 92% of patients. Gram negative organisms such as pseudomonas, klebsiella pneumonia, E.coli, Aerobacteraerogenes, Proteus, and Salmonella are the next most common organism. As anaerobic culture techniques improve, these organisms are recognized with increasing frequency.

Pathogenesis:
Empyema results from secondary to blockade of pulmonary lymphatics by inflammatory debris or the direct extension of the pneumonic process into the pleural space.
The American thoracic society (1962) has classified empyema into three phases based on the natural history of disease:

1. Exudative or acute phase- characterized by a pleural fluid of low viscosity with a cellular content. During this phase, the lung is still expandable.
2. Fibrin purulent or transitional phase- characterized by appearance of more turbid fluid due to an increase in polymorph neutrophils. Fibrin is deposited on both pleural surfaces, forming a limiting peel that prevents extension of empyema but also begins to trap and fix the lung. The pleural fluid becomes increasingly more turbid and the lung progressively less expandable with time.
3. Organizing or chronic phase- characterized by organization of the pleural peel, with ingrowth of capillaries and fibroblasts. The pleural fluid is very viscous, consisting of 75% sediment on standing. The organization can begin as early as 7 to 10 days after the onset of the disease, and usually by 4 to 6 weeks the process has entered the chronic phase.

III. Diagnosis:
Diagnosis of loculated pleural effusion/empyema is primarily made on the basis of Clinical features; Imaging, Pleural fluid examination and bronchoscopy.
Clinical features are not specific and Common symptoms include– pleuritic chest pain, Fever, Cough with sputum production, tachypnea, tachycardia, decreased respiratory excursions, pain on percussion, friction rub and absent breath sounds.
Radiographic examination should include anteroposterior and lateral chest radiographs which demonstrate parenchymal infiltrates or consolidation and pleural space fluid. CT and ultrasonography can delineate the nature and degree of parenchymal disease and character of the pleural fluid or rind when complete pacification of hemi-thorax is noted on plain films.
Bronchoscopy should be done to rule out endobrochial lesions such as tumors, inhaled foreign body or bronchopleural fistula.
In acute stage thoracocentesis sample should reveal, one or more of following features.

ph<7.2
Glucose < 40 mg/dl
LDH > 1000 IU/dl
Protein > 2.5
WBC > 500 /µ
Specific gravity > 1.018

Thin serous or cloudy fluid generally sterile.

In fibrin purulent or intermediate stage the fluid becomes thicker, opaque or fluid with positive cultures and in organizing or late stages an organizing peel with entrapment of the lung is seen. A sedimentation fraction of 75% indicates presence of thick walled empyema and open drainage is safe.

**IV. Management:**

The principals of management have been recognized by Hippocrates and ancient physicians of Greece10 and involved aspiration for diagnosis, repeated aspirations if warranted and tube or open drainage procedures once the cavity was stable.

**The Objectives Of Therapy Are:**

- Control of local and systemic infection with specific antibiotic therapy.
- Evacuation of empyema by one of a variety of methods selected on the basis of the etiology and stage of empyema process.
- Re-expansion of the lung with obliteration of the pleural dead space.

Needle thoracocentesis for chemistry analysis and culture is usually the initial diagnostic (and occasionally therapeutic) step coincident with the initiation of intravenous antibiotics. Thin exudates can occasionally be completely evacuated with this maneuver. Thoracocentesis and antibiotics alone have been successful in treatment of empyema in 6 to 20% of patients, particularly with early stage disease.11, 12, 13, 14 Large closed-tube thoracotomy with or without the adjunctive use of fibrinolytic agents has been traditional management with the fibrin purulent stage of the empyema process with reported success rates of 24 to 78%. The organizing phase of empyema requires direct removal of the restrictive coagulum (decortication) with open or laparoscopic technique.

**Empyectomy**

This refers to a specific manoeuvre - the complete excision of a small empyema without spillage of pus. A previous drainage procedure or the presence of a broncho-pleural fistula precludes this operation. Operative details have been well described by Dugan and Samson. Dissection is begun in the extra pleural plane and when the edges of the empyema sac are reached, the surgeon 'turns the corner' and then decorticates the inner surface of the empyema from the visceral pleura. Dissection is particularly difficult over the diaphragm where the endo thoracic fascia is thin and the extra pleural plane poorly developed.15

**Microvascular free muscle flaps**

There are patients in whom multiple previous operations to deal with a chronic empyema, usually with a broncho-pleural fistula, have failed. Under normal circumstances the use of a muscle flap would be considered but if the local muscles have been divided at previous thoracotomies, these muscles are frequently small and atrophied. In these circumstances Chen et al. have used micro vascular free muscle flaps. Contralateral latissimus dorsi and the lower four digitations of serratus anterior are mobilized on the same vascular pedicle, namely the thoracodorsal artery and vein. These muscle flaps are then anastomosed to the remnants of the thoracodorsal artery and vein on the affected side. He states these vessels are easily found. His results in his small series are excellent. The fistula is closed and the empyema space is obliterated. Or unable to cope with a permanent open drain, in whom decortication is undesirable or unsuccessful, is the procedure initially described by Clagett and Geracim for management of postpneumonectomy empyema. The same principles may be used for patients with an empyema who have had a limited or no previous pulmonary resection.16,17,18

**Thoracoplasty**

The purpose of thoracoplasty is to remove the rigidity of the outer chest wall and so establish contact between the now flexible chest wall and either residual lung; or, after pneumonectomy, with the mediastinum in order to obliterate the empyema space. The procedure is rarely undertaken today because of the deformity that results and the success of other forms of management.

**V. Material and methods;**

This study was conducted in department of CVTS in Sher I Kashmir institute of medical sciences soura. The study was a prospective randomized controlled analysis of all cases of empyema from Sep. 2011 to oct. 2013. This was a prospective randomized comparative case/ control study, between radiologically assisted
chest tube drainage and thoracotomy for loculated pleural effusion/empyema. Study was comprised of patients treated with radiographic chest tube drainage with the aid of streptokinase and patients treated with open thoracotomy. Both the groups were randomized using randomization table before a particular form of treatment was instituted.

The patients were evaluated in detail. Detailed history was taken from patients (or attendants) and both general physical examination as well as systemic examination was performed. Complete haemogram, serum chemistry, chest radiograph, abdominal ultrasonography and contrast enhanced computed tomography were performed.

**Inclusion criteria:** patients of all age groups, both sexes, patients giving informal written valid consent for the procedure, patients having radiologically documented loculated pleural effusion/empyema, patients having supportive / confirmative evidences of empyema from analysis of the thoracoacentrically aspirated fluid.

**Exclusion criteria:** patients not giving valid consent, patients with recent history of septicemia, patients on corticosteroids or immunosuppressents, patients needing surgical intervention for concurrent lung/thoracic lesions., patients having loculative effusion/empyema at more than two sites.

The method used in our study for drainage of loculative pleural effusion/empyema was a guided approach that is chest tube was inserted under ultrasound guidance and drainage observed for 24-48 hours and radiograph done to look for resolution. If radiograph revealed same findings what was observed before procedure then decision was taken to instill streptokinase. Following streptokinase instillation, drainage and radiographs were monitored for 7-10 days. If improvement did not occur then patient was treated with decortication. 25000iu of streptokinase were instilled in adults and in children 1500 IU/kg body weight dissolved in 50-100 ml of normal saline were used. Tube was clamped for 2-4 hours. Continuous data was statistically analysed by using two sample independent t test and paired t test. Data were analysed by using chi square/ fisher t test. P value <0.05 were considered statistically significant.

**Table showing comparison between cases and controls**

<table>
<thead>
<tr>
<th>Subjects</th>
<th>Radiographic assisted ICTD +Fibrinolytic</th>
<th>Decortication</th>
<th>P-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Length of hospital stay</td>
<td>13.6±2.362</td>
<td>7.9±1.837</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Average drainage</td>
<td>1387.5±477.27</td>
<td>703.57±159.48</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>ICTD removal(in Days)</td>
<td>15.7±1.9</td>
<td>7.9±1.6</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Pain after procedure</td>
<td>Discomfort related to chest tube</td>
<td>Discomfort related to chest tube and thoracotomy</td>
<td></td>
</tr>
<tr>
<td>Complete radiographic resolution</td>
<td>26</td>
<td>38</td>
<td></td>
</tr>
<tr>
<td>Success</td>
<td>26/40</td>
<td>39/40</td>
<td></td>
</tr>
<tr>
<td>Total cost procedure in rupees</td>
<td>8300±1316.6</td>
<td>610±213.18</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Complications associated</td>
<td>Allergy(sometimes fatal), bleeding, Pain, bleeding, air leak</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hospital turnover</td>
<td>Decreased</td>
<td>Increased</td>
<td></td>
</tr>
</tbody>
</table>

**VI. Discussion:**

Thoracic empyema still represents a significant cause of morbidity, prolonged hospitalization and mortality up to 10%. Important and common cause of empyema include pulmonary infection, surgical procedures, trauma, spontaneous pneumothorax, sub diaphragmatic infection and esophageal perforation. Approximately 50% of all bacterial pneumonias develop in pleural effusion, and in 1/3 rd. of these patients the pleural effusion becomes organized, as were have seen in our study that 35% patients did not responded to guide ICTD+ streptokinase. The propensity for developing empyema varies considerably with the type of bacteria producing the common pneumonia, the setting in which the infection is acquired and the alteration in these produce by antibiotic therapy. The second most frequent cause of empyema is post-surgical empyema which accounts for 20% of all cases. In our case series the bacteriological association was found to be insignificant(P- value> 0.10) but tubercle bacilli was found to be a measure etiological agent in 35% of patient population. The clinical picture of thoracic empyema may also be seen with pulmonary embolism, acute pancreatitis, Dressler’s syndrome and tuberculosis. The secondary complications of the empyema process include broncho-pleural fistula, empyma necessitans and osteomyelitis of the ribs or spine, invasion of the mediastinum with pulmonary esophageal fistula or pericarditis and brain abscess. In our case series such complication were not observed, suggesting that our interventions were conducted at an earlier stage. Once the diagnosis of empyema thoracis have been made by thoracentesis and approximate gram stain and/ or culture, computed tomographic( CT) scan is used universally to identify underlying parenchymal disease, to distinguish...
empyema from lung abscesses and to very the presence of loculations, the latter being an important factor in treatment planning. Effective management of empyema requires control of infection by approximate antibiotic therapy, evacuation of pus and obliteration of the empyema activity. In our case series we utilized USG for detection and confirmation of loculations in most of the cases and in few patients the CT scan was utilized. The most important aspect of management is the prompt initiation of effective drainage. Closed chest tube drainage is the usual first step which may take a no. of forms including image guided catheter insertion. Patients frequently come to medical attention when the pleural fluid is not free flowing and closed drainage is unlikely to be successful. The options available to manage inadequate drainage include additional chest tubes, intrapleural fibrinolytics, VATS and thoracotomy decortication. The choice of additional modality drainage depends upon the presence of ongoing pleural sepsis, maturity of empyema, degree of restriction of lung function, familiarity with treatment modalities and debility of patient.

Bibliography

[7]. Thoumani V, Brady KM, Mansour KA, Mille J, Lee RB.
[17] Chen H, Tang Y, NoodhoffMS, Chang C. Microvascular free muscle flaps for chronic empyema with bronchopleural fistula when the major local muscles have been divided - one-stage resection with primary wound closure. Annals of Plastic Surg 1990;24:510-6