

Intrapleural Fibrinolytic Therapy in Loculated Effusion in a Pediatric Patient



Dr. Rhea Gandhi

Department of Respiratory Medicine





10 year old girl, k/c/o epilepsy on treatment, Weight: 29.8 kg (- 1 SD)

Chief Complaints:

- Fever 2 weeks
- Dry cough 2 weeks
- Loss of appetite 2 weeks
- Breathlessness (MMRC grade 2) 1 week
- Vomiting 5 days

No significant past history

No history of close contact with tuberculosis patient





Clinical Examination

- GCS-15/15
- Temp- 98.4° F
- PR-130 beats/ min
- RR- 22 breaths/ min
- **BP-** 104/72 mmHg
- SpO2-96% on room air

Admitted in Pediatric ward on 25/12/23

- Respiratory System- Reduced breath sounds on the left side of chest
- Other systems- Normal



INVESTIGATIONS



Investigation	Value	Investigation	Value
Hb	11.6 gm/dl	Total protein	7.2 g/dl
TLC	7600 /µl	S. LDH	218 U/Lt
N / L	57 / 32	CRP	77.90 mg/L
Platelet count	460000 /µl	ESR	48 mm/hr
Peripheral blood smear	Mild hypochromia, microcytosis & anisocytosis	Urine R/M	Pus cells 1-2 No RBCs
Bilirubin (T/ C/ U)	(0.21/ 0.10/ 0.11)	Mantoux test	Negative
SGOT/ SGPT/ ALP	23/12/183	Rapid malarial antigen	Negative
Urea/ Creatinine	21/0.43	Dengue profile	Negative
HIV	Negative	Widal	Negative



CLINICAL COURSE





USC thorax

- Left sided pleural effusion
- Approx 1000 1200 cc volume
- Incomplete internal septations
- Consolidation and collapse of underlying lung parenchyma







Started on antibiotics- Ceftriaxone, Vancomycin

Left sided Intercostal drainage tube was established on 25/12/23 in PICU Drain ~ 900 ml





PLEURAL FLUID ANALYSIS



Pleural fluid	Result	
Protein	5.7 gm %	
Glucose	81 mg/dl	
TLC	1600 per cmm	
ADA	47.40 U/L	
LDH	450 U/L	
Culture	No growth	
Staining (Gram, ZN)	Negative	
Gene Xpert	Mtb not detected	
Malignant cytology	No malignant cells seen	

Exudative lymphocytic effusion of Tubercular etiology







<image>

After ~ 1500 ml drain, there was no drain from intercostal tube over the next 4 days





CLINICAL COURSE



<u>Pediatric surgery consult was sought:</u> Advised Video assisted thoracoscopic surgery (VATS); SOS thoracotomy

Simultaneously, <u>Respiratory medicine</u> <u>consultation</u> was given for further management of pleural effusion. It was opined that the patient may not require VATS/ thoracotomy.

Transferred to female pulmonary ward for further management on 3/1/24







- Patient was still having fever spikes in spite of receiving 9 days of antibiotics
- ≻Repeat TLC was 7100 /µl
- >Antibiotics were stopped

>Patient was registered on DOTS

HRZ	E
50/75/150	100
3 + 1A*	3

- A* Adult FDC (HRZE = 75/150/400/275)
- H- Isoniazid
- **R-**Rifampicin
- Z-Pyrazinamide
- E-Ethambutol



COURSE IN RESPIRATORY MEDICINE



• USG thorax showed approx. 200 to 300 cc effusion with internal septations.

• In view of left loculated tubercular pleural effusion, it was decided to initiate Intrapleural Fibrinolytic Therapy for the patient.





PRE-REQUISITES OF IPFT

- IPFT is indicated only in those cases where significant loculated pleural fluid is present.
- > The tube or catheter should be correctly positioned and patent.
- The drainage from intercostal tube or catheter should be less than 50ml per day.











Age group	Streptokinase	Urokinase
6-12 years	l lakh IU	50000 IU
1-6 years	50000 IU	25000 IU
< 1 year	25000 IU	10000 IU

M S Barthwal. Intrapleural Fibrinolytic Therapy in Loculated Pleural Effusions. Journal of The Association of Physicians of India.Vol. 68. June 2020



CLINICAL COURSE







PRE IPFT



POST IPFT





DE D.Y. PATIL VIDYAPEETH, PUNE OPTIONS AFTER FAILED TUBE DRAINAGE





IPFT is a cost effective solution for draining loculated parapneumonic effusions and this option must be exercised in our country for eligible patients before considering surgery.



RATIONALE OF IPFT









RATIONALE OF IPFT





IPFT VS VATS



IPFT	VATS	
No anesthesia	Under GA, Single lung ventilation	
More accessible	Lesser accessibility	
Expertise not required	Expertise required	
Lesser invasive (ICD)	More invasive (2 or 3 ports)	
Cost effective	Significantly costlier	
Fibrinolysis	Adhesiolysis, debridement, decortication	
Less effective	More effective	



Safety and Efficacy of Streptokinase in Multiloculated Pleural Effusion in Pediatric Population



Akhtar Reza, Mohd Aslam, Manju Gupta¹, Mohd Azam Haseen², Mayank Yadav²

Departments of General Surgery and ²Cardiothoracic Surgery, JN Medical College, AMU, Aligarh, Uttar Pradesh, ¹Department of Cardiothoracic and Vascular Surgery, VMMC and Safdarjung Hospital, New Delhi, India

> Chest. 1993 Apr;103(4):1190-3. doi: 10.1378/chest.103.4.1190.

Intrapleural streptokinase as adjunctive treatment for persistent empyema in pediatric patients

H Rosen¹, V Nadkarni, M Theroux, R Padman, J Klein

ORIGINAL ARTICLE

Randomised trial of intrapleural urokinase in the treatment of childhood empyema

A H Thomson, J Hull, M R Kumar, C Wallis, I M Balfour Lynn, on behalf of the British Paediatric Respiratory Society Empyema Study Group

Thorax 2002;57:343-347



Intrapleural Streptokinase in a Two-Year-Old Child with a Parapneumonic Effusion



M.S. Barthwal

Department of Medicine, Base Hospital, Delhi Cantt, Delhi, India

ABSTRACT

A two-year-old child was hospitalised with features of parapneumonic effusion. He was initially managed with parenteral antibiotics and chest tube drainage. After three days drainage became insignificant inspite of chest tube being patent and appropriately positioned. CT scan of chest showed multiloculated effusion. In view of multiloculated effusion it was decided to try intrapleural fibrinolysis with streptokinase. Streptokinase in a dose of 1,25000 IU dissolved in 50 ml of normal saline was instilled through the chest tube daily. After instilling three doses, there was a significant increase in the drainage followed by almost complete radiological resolution. There were no side effects. Intrapleural streptokinase is a useful adjunctive threapeutic modality in the management of complicated parapneumonic effusion or empyema in paediatric patients.

Key words : Parapneumonic effusion, Streptokinase, Fibrinolysis.





M.S. Barthwal¹, V. Marwah², M. Chopra³, Y. Garg², R. Tyagi⁴, K. Kishore⁵, A. Vijay⁶, V. Dutta⁷, C.D.S. Katoch³, S. Singh⁵ and D. Bhattacharya¹

Department of Pulmonary Medicine, Military Hospital (Cardiothoracic Centre), Armed Forces Medical College¹, Pune; Army Hospital (R&R)², New Delhi; Military Hospital³, Ranchi; Indian Naval Hospital⁴, Mumbai; Command Hospital⁵, Lucknow; Command Hospital⁶, Chandimandir and Base Hospital⁷, Delhi Cantt, India

	No. (%)	Mean Age±SE
Gender		
Male	183 (91.5%)	
Female	17 (8.5%)	
Age (in years)		
Above 12	185 (92.5%)	31.7 ± 8.4
Below 12	15 (7.5%)	6.4±2.4
Aetiology of pleural effusio	n	
CPE	106 (53%)	
Tubercular	59 (29.5%)	
Empyema	23 (11.5%)	
Traumatic hemothorax	12 (6%)	

Definition of abbreviations: SD=Standard deviation; CPE=Complicated parapneumonic effusions Response rate to Streptokinase and Urokinase was similar.

Indian J Chest Dis Allied Sci. 2016

> The interval between the onset of loculated pleural collection and initiation of IPFT was more than 6 weeks in non responders.

Adverse effects observed: mild chest pain, low-grade transient fever.







Intrapleural Fibrinolytic Therapy in Loculated Pleural Effusions

MS Barthwal

Journal of The Association of Physicians of India • Vol. 68 • June 2020

Abstract

About 36% to 57% of bacterial pneumonias develop parapneumonic effusion. When the chest tube is correctly positioned as evidenced by postero-anterior and lateral chest radiographs and there is a significant amount of pleural fluid, the major reasons for failed drainage are multiple pleural space loculations or tube obstruction by thick and viscous fluid. The various modalities of treatment available for loculated pleural effusion are: saline flushes, placing one or more catheters in loculi under image guidance, video assisted thoracoscopic surgery (VATS), standard thoracotomy with drainage of empyema and decortication. The first two modalities are not so effective in improving drainage. The last two surgical modalities are more invasive, not easily available and, if available, are not affordable by majority of patients in the developing countries like India. The fibrinolytic agents, if used early in loculated pleural effusions, break loculations and early pleural peel thereby facilitating pleural space drainage.

1949 in 23 patients who had loculated empyema or hemothorax. Their patients received intrapleural instillation of both streptokinase and streptodornase, which was extracted from concentrated filtrates of streptococci of Lancefield group C. There was significant improvement in drainage of pleural fluid. However, the initial enthusiasm waned because of significant systemic adverse effects in the form of fever, leukocytosis and general malaise. These side effects were due to immunological reaction caused by impurities in the preparation of agents. There was not much of use of this therapy until Bergh and colleagues4 in 1977 used purified





TAKE HOME MESSAGE

 ✓ IPFT is a safe and cost effective option in the management of loculated effusions of varied etiologies and also in <u>pediatric patients</u>.

 For a developing country like ours, this option must be exercised in eligible patients before subjecting them to costlier, not so easily accessible and more invasive surgical options.





