

Recombinant Factor VII

Sir,

The paper by Teh *et al*¹ stated that recombinant factor VII_a was used for the first time in the region for a patient with acquired haemophilia at University Hospital, Kuala Lumpur. For the record, the first use of factor VII_a in an acquired haemophilia in Malaysia was for a 26-year-old man with life-threatening retroperitoneal haemorrhage at Subang Jaya Medical Centre in March 1992. The case was presented as an oral paper in the first National Conference of Haematology in April 1994 and published as a case report² in *Annals of Academy of Medicine (Singapore)* in 1994.

References

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Chylothorax 2° to Tuberculosis

Sir,

Chylothorax is a rare condition. Tumour accounts for about 50% of cases. Tuberculosis is not a common cause of chylothorax. We report a case of chylothorax

in which all the other common causes were ruled out and which responded to anti-tuberculous therapy.

A 40-year-old Chinese lady presented with a recent history of non tender left neck swelling, associated with low grade fever, night sweats and mild dyspnoea on exertion. There was no history of tuberculosis or contact with tuberculous patients. Clinically there was some fullness on the left side of the neck and findings consistent with a left pleural effusion.

Investigations revealed a raised ESR at 80 mm/hour and 150 mm/hour on 2 consecutive examinations. Mantoux test was positive at 16 mm. A chest X-ray confirmed a left pleural effusion. Pleural tap revealed milky fluid which was alkaline (pH 7.92) and consistent with exudate (Albumin 32 gm/L). Lipoprotein electrophoresis showed presence of chylomicrons and VLDL. The Sudan III stain was positive.

Pleural Fluid

Total Cholesterol (TC) : 7.25 mmol/L
Triglycerides (Tg) : 29.00 mmol/L

Fasting Serum Lipids

Total Cholesterol (TC) : 4.32 mm/L
Triglycerides (Tg) : 1.10 mm/L

The IFAT (Indirect Fluorescent Antibody Technique) for filariasis showed normal titres. Bronchoscopy was normal. She was started on a trial of anti tuberculous treatment (2HR²/4RH) and discharged one week later. At follow-up after two weeks, she had CT scan Thorax/Abdomen which was normal. A repeat CXR done two weeks later showed the effusion had cleared. She was then put on a full 6 months course of treatment and is presently doing well on follow-up.

The only other condition which may present similarly is pseudo-chylothorax. Some of the points in

differentiating between chylothorax and pseudo-chylothorax are as follows¹:

	Chylothorax	Pseudo-Chylothorax
Presentation	More acute	Always chronic
Pleural surface	Smooth	Thickened/ calcified
Cause of opalescence	Chylomicrons	Cholesterol crystals or lecithin-globulin complexes
Triglyceride levels	High ($> 110 \text{ mg dl}^{-1}$)	Usually low
Cholesterol	Low	Usually high
Chylomicron on lipo-protein electrophoresis	Yes	No

We feel that our patient had chylothorax as the presentation was rather acute, the triglyceride levels in the pleural fluid was very high despite a normal serum triglyceride level and there was presence of chylomicrons on lipoprotein electrophoresis with a positive Sudan III stain.

The mechanism by which TB causes chylothorax is unclear although a possibility is the extrinsic compression of the thoracic duct by inflamed nodes². The CT scan (Thorax) in our patient was normal but it may have missed the enlarged nodes as it was done 3 weeks after anti TB-treatment (by which time the nodes may have regressed).

Finally, this case illustrates that although tuberculosis is common in our country, it can still surprise us with atypical presentations.

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2. Bessone LN, Ferguson TB, Burford TH. Chylothorax Ann Thorax Surgery 1971;12 : 527-50.