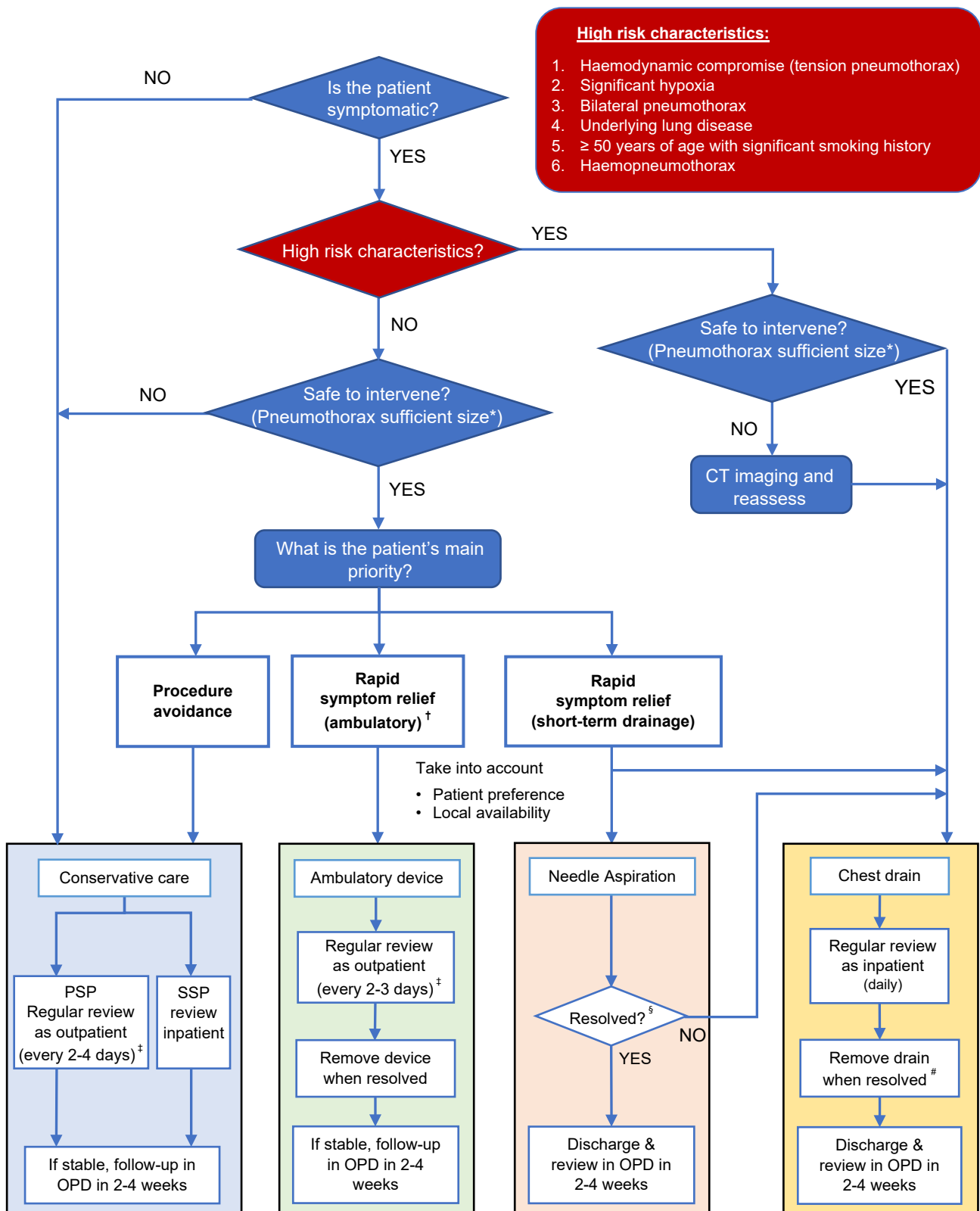


# Appendix 1 – Clinical pathways/decision trees

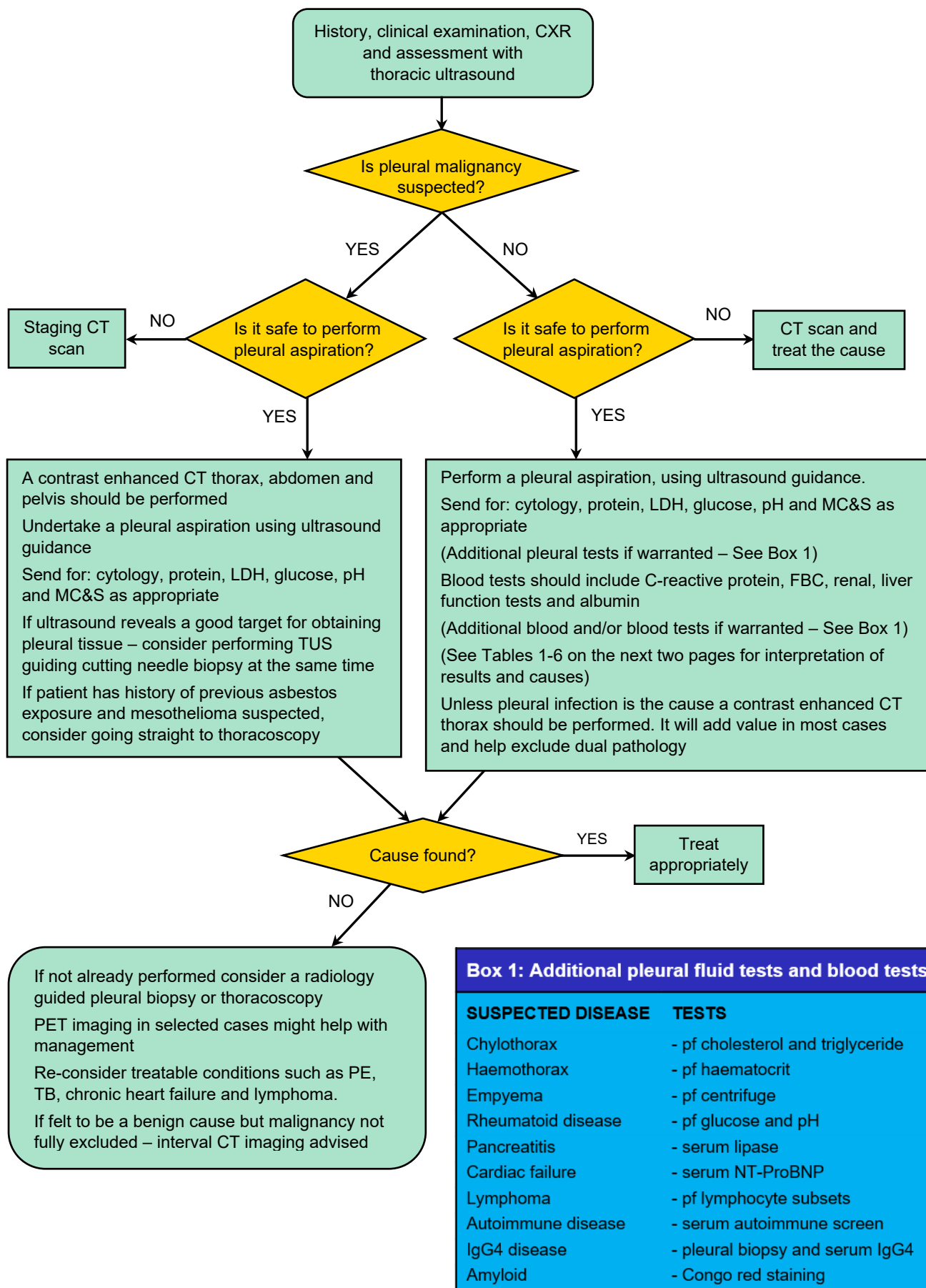
## Pneumothorax Pathway



\* Pneumothorax of sufficient size to intervene depends on clinical context but, in general, usually ≥ 2cm laterally or apically on CXR, or any size on CT scan which can be safely accessed with radiological support.  
<sup>†</sup> If ambulatory pathway available locally.  
<sup>‡</sup> At review, if enlarging pneumothorax or symptoms consider chest drain insertion and admission.  
<sup>§</sup> Success: improvement in symptoms and sustained improvement on CXR.  
<sup>#</sup> Talc pleurodesis can be considered on the first episode of pneumothorax in high risk patients in whom repeat pneumothorax would be hazardous (eg, severe COPD).

CXR, chest X-ray; COPD, chronic obstructive pulmonary disease; OPD, outpatient department; PSP, primary spontaneous pneumothorax; SSP, secondary spontaneous pneumothorax.

# Unilateral pleural effusion diagnostic pathway



CXR, chest X-ray; FBC, full blood count; LDH, lactate dehydrogenase; NT-proBNP, N-terminal prohormone brain natriuretic peptide; PE, pulmonary embolism; TB, tuberculosis; TUS, thoracic ultrasound.

## Unilateral pleural effusion diagnostic pathway – Tables 1-3

**Table 1**

<b>Light's criteria</b>
Pleural fluid is an exudate if one or more of the following criteria are met:
<ul style="list-style-type: none"> <li>• Pleural fluid protein divided by serum protein is <math>&gt;0.5</math></li> <li>• Pleural fluid lactate dehydrogenase (LDH) divided by serum LDH is <math>&gt;0.6</math></li> <li>• Pleural fluid LDH <math>&gt;2/3</math> the upper limits of laboratory normal value for serum LDH</li> </ul>

**Table 2**

<b>Transudates</b>	<b>Exudates</b>
<p><b>Common</b></p> <ul style="list-style-type: none"> <li>• Congestive cardiac failure</li> <li>• Liver cirrhosis</li> <li>• Hypoalbuminaemia</li> <li>• Nephrotic syndrome</li> </ul>	<p><b>Common</b></p> <ul style="list-style-type: none"> <li>• Malignancy</li> <li>• Pleural infection</li> <li>• Pulmonary embolism</li> <li>• Autoimmune pleuritis</li> </ul>
<p><b>Less common</b></p> <ul style="list-style-type: none"> <li>• Nephrotic syndrome</li> <li>• Mitral stenosis</li> <li>• Peritoneal dialysis</li> <li>• Chronic hypothyroidism</li> <li>• Constrictive pericarditis</li> </ul>	<p><b>Less common</b></p> <ul style="list-style-type: none"> <li>• Drugs</li> <li>• Lymphatic disorders</li> <li>• Meigs syndrome</li> <li>• Post-coronary artery bypass graft</li> <li>• Benign asbestos related pleural effusion</li> </ul>

**Table 3**

<b>Causes of lymphocytic pleural effusion</b>
Malignancy
Tuberculosis
Lymphoma
Congestive cardiac failure
Post-coronary bypass graft
Rheumatoid arthritis
Chylothorax
Yellow nail syndrome

## Unilateral pleural effusion diagnostic pathway – Tables 4-6

**Table 4**

Causes of bilateral pleural effusions
Congestive cardiac failure
Hypoalbuminaemia
Renal failure
Liver failure
SLE and other autoimmune diseases
Widespread malignancy including abdominal/pelvic malignancy
Bilateral pulmonary embolus

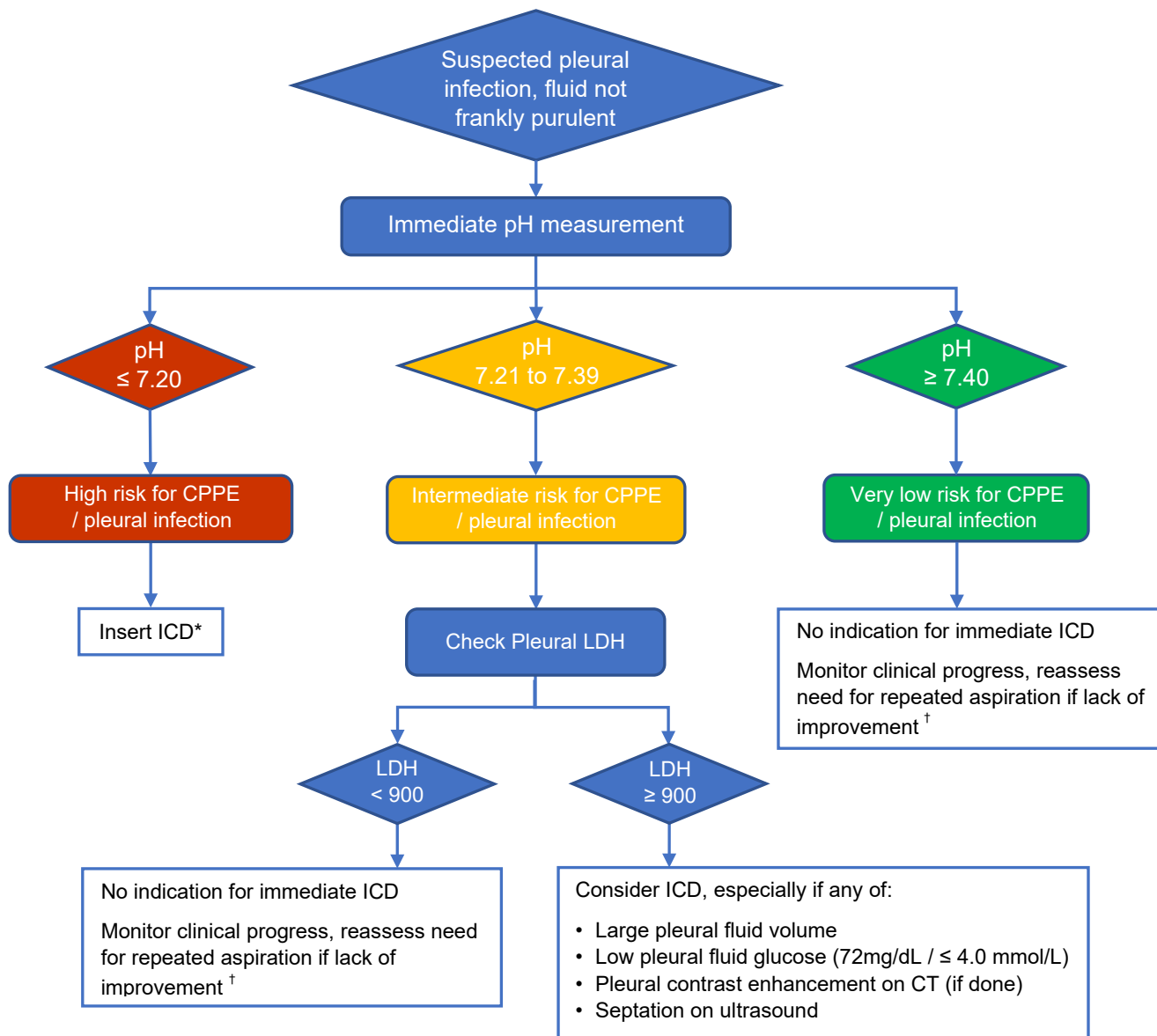
**Table 5**

Pleural fluid lipid values in chylothorax and pseudochylothorax	
<b>Chylothorax:</b>	
• Triglycerides	– high >1.24 mmol/L (110 mg/dL)
• Cholesterol	– low
• Cholesterol crystals	– absent
• Chylomicrons	– usually present
<b>Pseudochylothorax:</b>	
• Triglycerides	– low
• Cholesterol	– high >5.18 mmol/L (200 mg/dL)
• Cholesterol crystals	– often present
• Chylomicrons	– absent

**Table 6**

Causes of chylothorax and pseudochylothorax	
<b>Chylothorax:</b>	
• Trauma:	thoracic surgery (especially if involving posterior mediastinum, eg, oesophagectomy), thoracic injuries
• Neoplasm:	lymphoma or metastatic carcinoma
• Miscellaneous:	disorders of lymphatics (including lymphangiomyomatosis), tuberculosis, cirrhosis, obstruction of the central veins, chyloascites
• Idiopathic (about 10%)	
<b>Pseudochylothorax:</b>	
• Tuberculosis	
• Rheumatoid arthritis	

# Suspected pleural infection, non-purulent fluid – initial decision tree

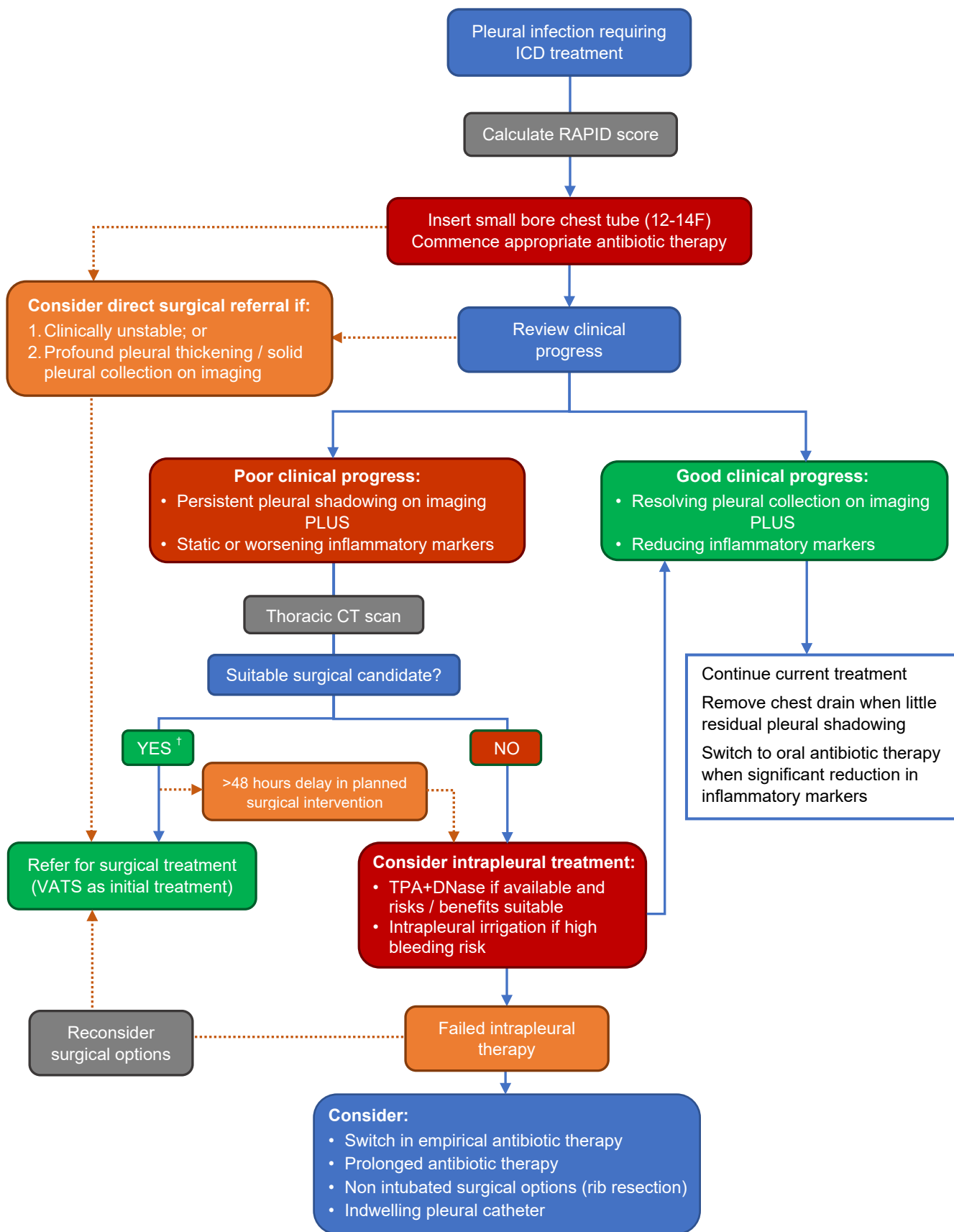


\* Assuming ultrasound demonstrates safe volume of accessible pleural fluid.  
 † As evidenced by ongoing temperature, persisting elevation of inflammatory markers. Those with septations and pleural pH >7.4 should also be considered for drainage.

Initial pH	Level of risk for CPPE / pleural infection	Initial action regarding drainage
≤ 7.2	High risk	Insert ICD, assuming ultrasound demonstrates safe volume of accessible pleural fluid
> 7.2 to < 7.4	Intermediate risk	Check LDH and review other parameters which may support CPPE / pleural infection. Consider ICD insertion if LDH > 900, especially if any of the following: <ul style="list-style-type: none"> <li>• Large pleural fluid volume</li> <li>• Low pleural fluid glucose (72 mg/dL / ≤ 4.0 mmol/L)</li> <li>• Pleural contrast enhancement on CT</li> <li>• Septation on ultrasound</li> </ul>
≥ 7.4	Very low risk	No indication for immediate ICD

CPPE, complex parapneumonic effusion; LDH, lactate dehydrogenase; ICD, intercostal drain.

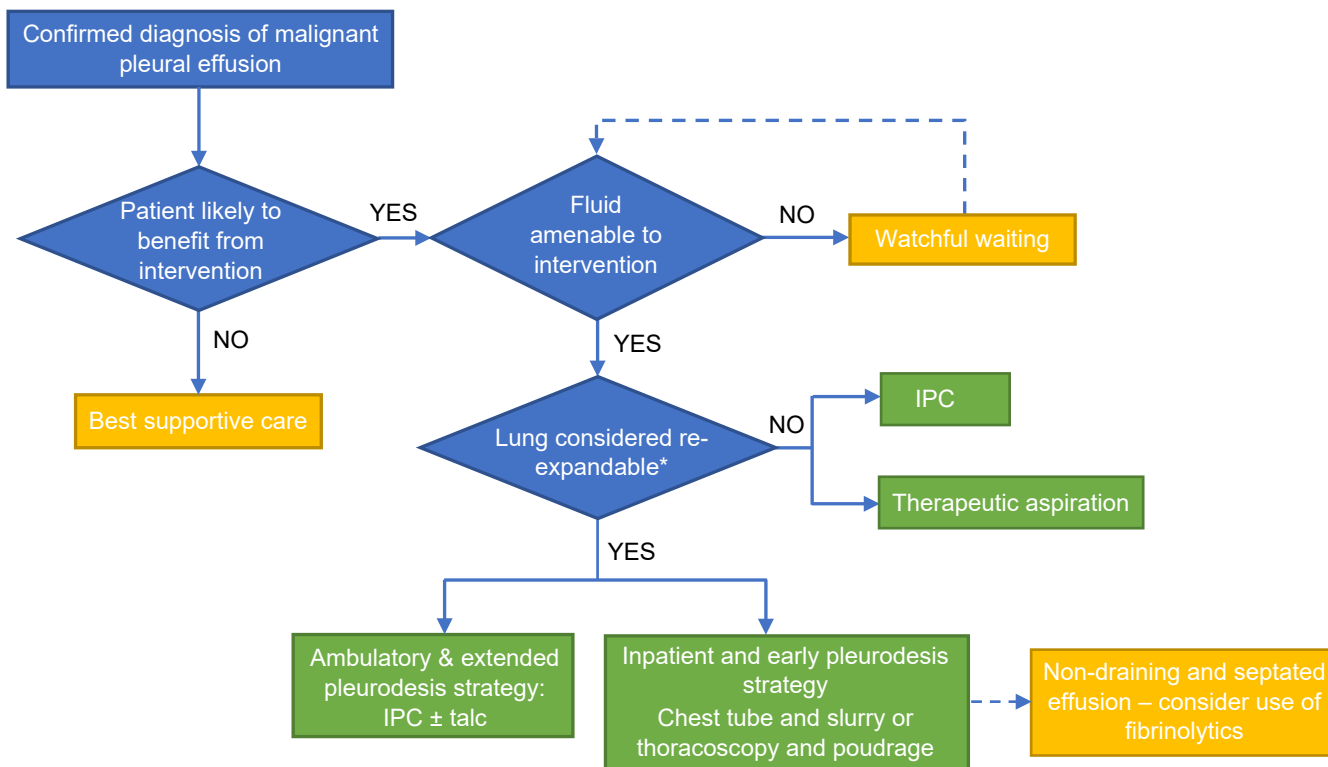
# Pleural infection treatment pathway



\* Antibiotic therapy should be based on likely organisms initially and adapted according to positive culture results, with consideration of anaerobic cover throughout.  
 † Intrapleural treatment may be considered prior to surgical treatment in liaison with surgical expertise.

ICD, intercostal drain; TPA, tissue plasminogen activator; VATS, video-assisted thoracoscopy surgery.

# Malignant pleural effusion pathway



\* Review of imaging, possible trial of benefit of aspiration before final decision.

IPC, indwelling pleural catheter.