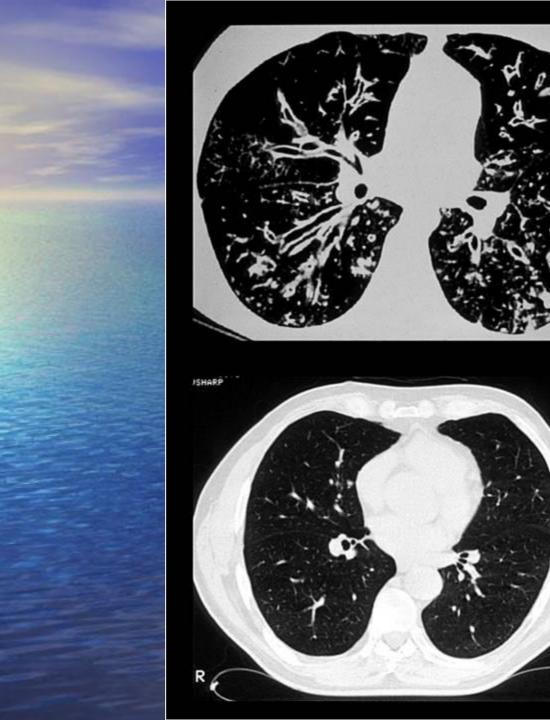
Bronchiectasis

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Bronchietasis

Originally described by Laennec in 1819 Chronic Debilitating Characterised persistent cough excessive sputum production recurrent chest infection

Permanent abnormal dilatation
impaired mucociliary clearance
bacterial colonisation
excessive airways inflammation





Bronchiectasis

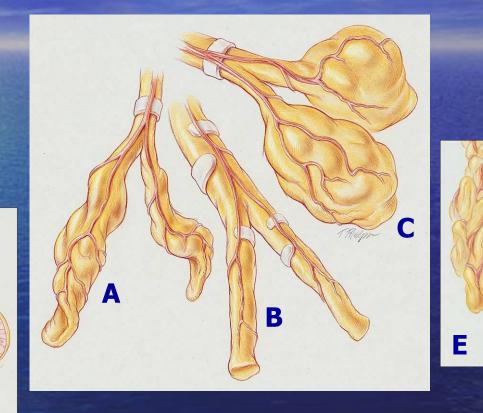
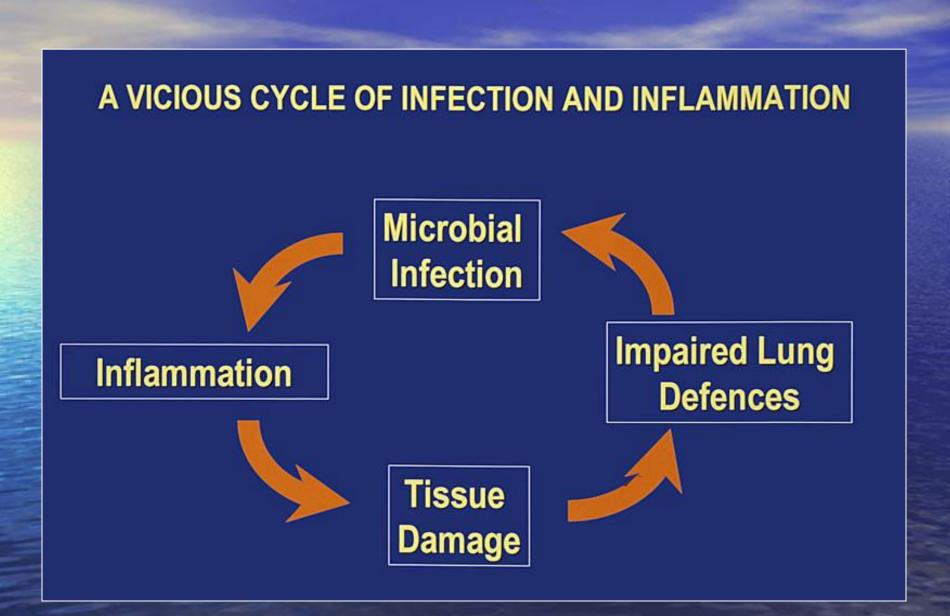


Figure 14–1. Bronchiectasis. A, Varicose bronchiectasis. B, Cylindrical bronchiectasis. C, Saccular bronchiectasis. Also illustrated are excessive bronchial secretions (D) and atelectasis (E), which are both common anatomic alterations of the lungs in this disease.

D



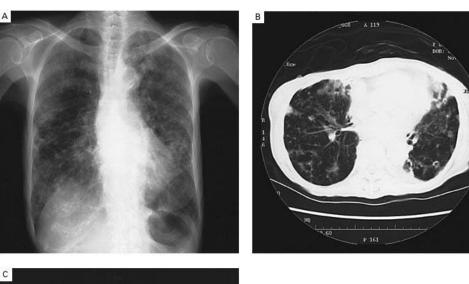
Etiology Acquired bronchiectasis Recurrent pulmonary infection Bronchial obstruction - Childhood infection e.g measles, pertussis Aspiration Congenital bronchiectasis Kartagener's syndrome - Hypogammaglobulinemia – Cystic fibrosis Abnormal cartilage formation

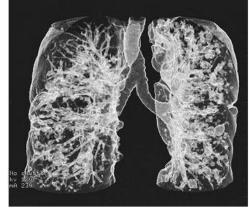
Aetiology of bronchiectasis

Cause	n (% of study)
Post infection	51 (32)
Idiopathic	42 (26)
PCD	17 (11)
ABPA	13 (8)
Immune deficiency	9 (6)
Ulcerative colitis	5 (3)
Young's syndrome	5 (3)
Pan bronchiolitis	4 (3)
Yellow nail syndrome	4 (3)
Mycobacterium infection	4 (3)
Rheumatoid arthritis	3 (2)
Aspiration	2 (1)
CF variant	2 (1)
Total	161

ABPA = Allergic bronchopulmonary aspergillosis, PCD = Primary ciliary dyskinesia, CF = Cystic fibrosis

An 81-year-old woman was admitted with weight loss (18 kg in 27 months), hemoptysis, and tubular and diffuse granular shadows on her chest radiograph (Panel A)

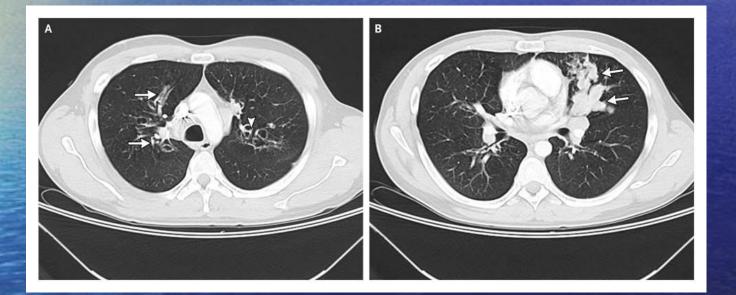




Final diagnosis: MAC infection of bronchiectasis

Ebihara T and Sasaki H. N Engl J Med 2002;346:1372

A 26-year-old man who smoked and had a long history of poorly controlled asthma and severe environmental allergies was admitted for an exacerbation of asthma

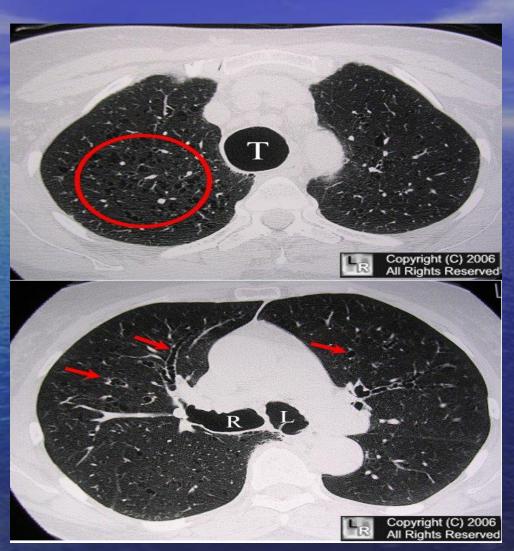


Final diagnosis: ABPA

Chen T and Hollingsworth H. N Engl J Med 2008;359:e7

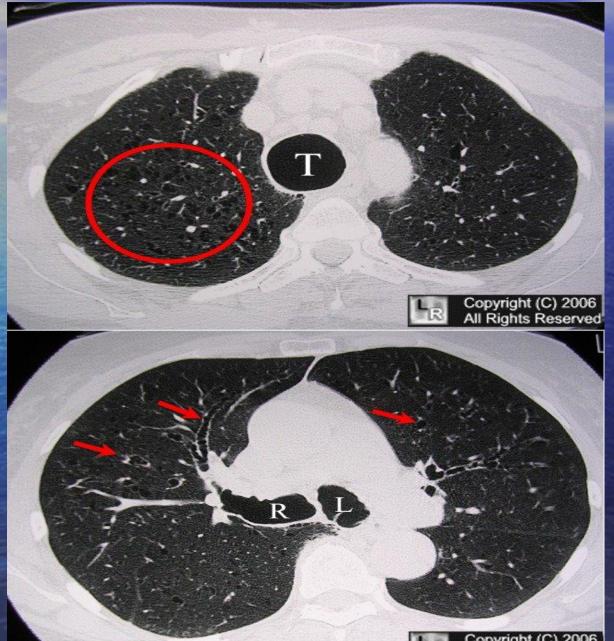
PCD Katergener's





Mounier-Kuhn P. Dilatation de la trachee: Constatations radiographiques et bronchoscopiques. Lyon Medical. 1932;150:106–9.

Mounier-kuhn



Bronchiectasis

Adults Who to suspect

Persistent productive cough

young age at presentation
symptoms over many years
absence of smoking history
daily expectoration of large volumes of sputum
haemoptysis

Unexplained

haemoptysis

non-productive cough

After excluding other causes

HISTORY WHICH SHOULD LEAD TO SUSPICION OF BRONCHIECTASIS

Recurrent LRTI Chronic productive cough Breathlessness, wheeze Haemoptysis Chest pain Tiredness (ENT, infertility, GI, ILD)

Thought to have COPD

COPD with Bronchiectasis no history of smoking • there is slow recovery from lower respiratory tract infections recurrent exacerbations Sputum growth/colonised with Pseudomonas aeruginosa

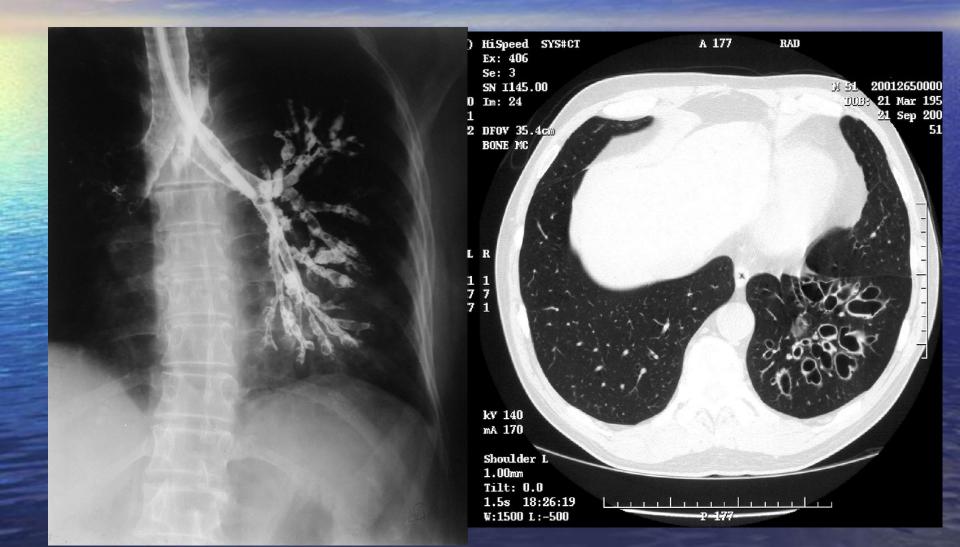
Investigations

CxrayHRCT

Sputum MCS
1. When stable
2. Onset exacerbating

Spirometry

Radiology



Exacerbations

Is it an exacerbation 0 **?Antibiotics required** 0 1. Deterioration over days **Increasing Cough** Increased sputum volume or change of viscosity 3. increased sputum purulence + increasing wheeze & 4. breathlessness 5. haemoptysis 6. systemic upset 7. Non specific

Antibiotic Choice, Dose and Duration

Admit

Development of cyanosis or confusion Breathlessness with a respiratory rate >25/minute Circulatory failure, respiratory failure, cyanosis or confusion Temperature >38°C Patient unable to take oral therapy Patient unable to cope at home Haemoptysis >25mls/day Intravenous therapy required in patients with clinical failure after oral antibiotics

Common organisms associated with acute exacerbation of bronchiectasis and suggested antimicrobial agents

Streptococcus pneumoniae Amoxicillin 500 mg tds Clarithromycin 500 mg bd 14 days

Haemophilus influenzae (b-lactamase negative)

Amoxicillin 500 mg tds Amoxicillin 1 g tds Amoxicillin 3 g bd Clarithromycin 500 mg bd

Haemophilus influenzae (b-lactamase positive)

Co-amoxiclav 625 mg tds Clarithromycin 500 mg bd Ciprofloxacin 500 mg bd

Moraxella catarrhalis Co-amoxiclav 625 mg tds Ciprofloxacin 500 mg bd

Staphylococcus aureus (MSSA) Flucloxacillin 500 mg qds Clarithromycin 500 mg bd

MRSA

Coliforms Ciprofloxacin

Pseudomonas

Empiric therapy

Amoxycillin 500mg tds 14days Clarithromycin 500 bd Severe Bronchiectasis/colonised with H influenzae Amoxycillin 1g tds/ 3g bd Pseudomonas colonised patients Ciprofloxacin 500/750 bd.

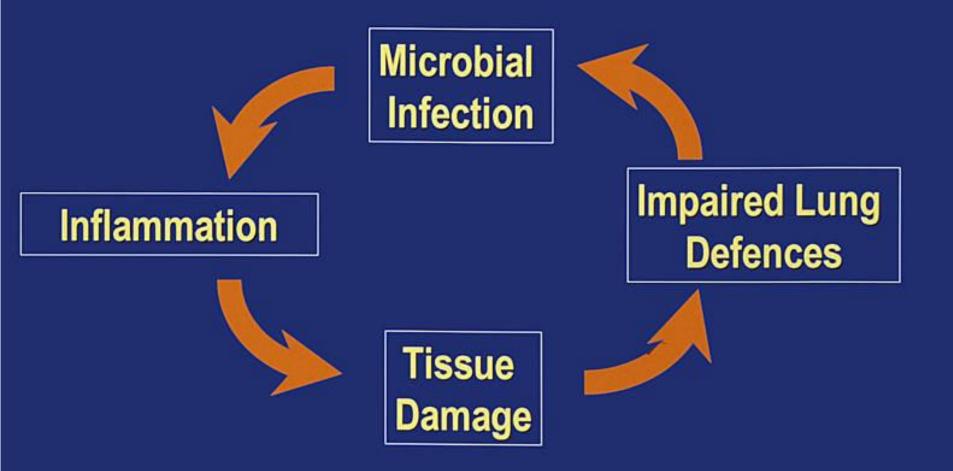
Long Term antibiotics

=>3 Exacerbations/yr
Fewer Exacerbation but significant morbidity
Nebulised antibiotics Gent/tobramycin/colistin
Long term Macrolides

Management

Physiotherapy
Immunisation
Bronchodilators
Mucolytics
Nebulised saline

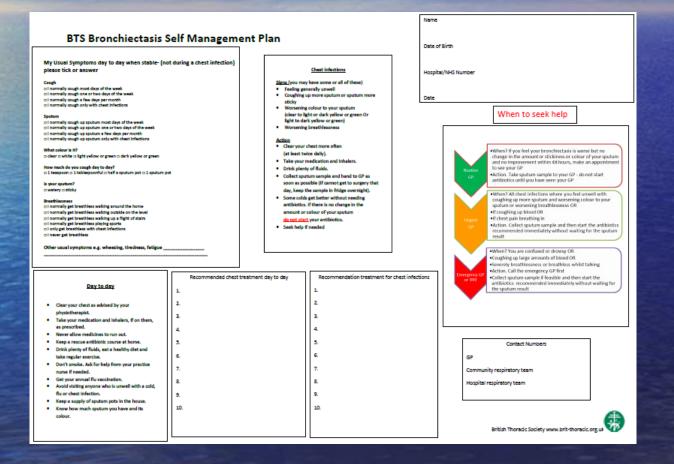
A VICIOUS CYCLE OF INFECTION AND INFLAMMATION



Monitoring

Symptom
Sputum Volume 24hrs/Purulence
Frequency of Exacerbations/yr
Frequency of Antibiotic use
FEV1 FVC PEF annually
Cxray only if indicated

Self Management Plan



A VICIOUS CYCLE OF INFECTION AND INFLAMMATION

