

()Prognosis:

- 1-age
- 2-post-bronchodilator FEV1
- 3- wt loss: <21 BMI
- 4- pulmonary hpt
- 5- degree of airway obstruction: FEV1<35
- 6- Excersize capacity: poor if < 150 m
- 7- measurment of dyspnea: if MRC > G4

()Acute exacerbation of COPD:

- *characterised by an increase in synptoms & deterioation in lung function.
- *caused by bacteria, viral & change in air quality
- *many pt can be managed at home with the use of increased dose of bronchodilators, steroid & AB.

So admission to hospital indicated in:

- 1- presence of cynosis
- 2- peripheral odema
- 3 presence of comorbidity
- 4- altered level of conc
- 5-Inadequate response of symptoms to outpatient management
- 6- Marked increase in dyspnea
- 7- Inability to eat or sleep due to symptoms
- 8- Worsening hypoxemia Worsening hypercapnia
- 9- Changes in mental status Inability to care for oneself (ie, lack of home support)
- 10- Uncertain diagnosis

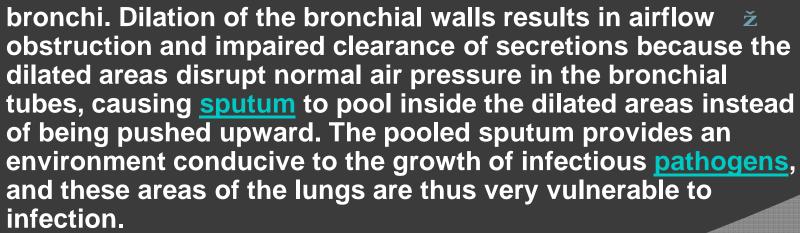
in hospital do the followings: ()check PaO2, PaCO2, PH, CXR, ECG, CBC, BU, electrolytes, PFT & sputum for culture & sensetivity. ()02 therapy: ()bronchodilators: ()oral CS: 30 mg for 10 days indicated in: 1- if pt already on oral CS 2- if prevoluos response of CS 3- failure to response to bronchodilators 4-1st presentaion of disease ()AB: ()diuretics ()heparin ()NIV

()Complications:

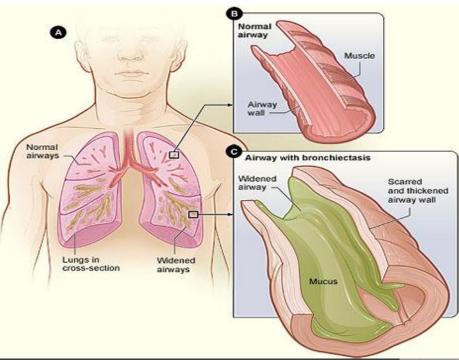
- *resp failure
- *core-pulmonale
- *rupture of bullae lead to pneumothorax
- *amylodosis

Bronchiectasis: ž

()definition: abnormal dilatation of



When this happens, the bronchial tubes become more inelastic and compressed, creating a self-perpetuating cycle of further damage to the lungs.



() types: ž

There are three types of brochiectasis, varying by level of severity.

- 1- Fusiform (cylindrical) bronchiectasis (the most common type) refers to mildly inflamed bronchi that fail to taper distally.
- 2- In varicose bronchiectasis, the bronchial walls appear beaded, because areas of dilation are mixed with areas of constriction.
- 3- Saccular (cystic) bronchiectasis is characterized by severe and irreversible ballooning of the bronchi peripherally, with or without airfluid levels.

()ateology:

1- congenital: *cystic fibrosis

*ciliary dysfunction syndroms include primary ciliary dyskinesia & kartagner's

*primary hypogammaglobulinemia ž

2- aquired-children: *pneumonia 🛛 🎽

* primary TB 👱

*inhaled foreign body ž

```
3- acquired-adults: *suppurative pn
                   *pulTB
                   *Allergic broncopulmonary aspergillosis
                   *bronchial tumor
Clinical features: ½
A- symtopms: \geq
*cough: ž
*pleurisy: ž
*hemoptysis: (dry bronchiectasis) 👱
*wt loss, anorxia, lassitude & digital clubbing. 🔀
B- signs: ž
*may be asymptomatic, either unilateral or bilateral. 
*crackles, wheezing. *\otimes
*later on bronchial breathing due to pul fibrosis. 🔀
```

()Investigations: •

- 1- sputum: ž
- 2- CXR: cystic bronchiactetic changes
- 3- CT scan: ž
- 4- assessement of ciliary dysfunction ž



()physiotherapy:

()AB: Oral ciprofloxacin 250-750 mg 12 h or ceftazidime 1-2 g

()surgery: young pt, unilateral, small area.

() Complications:

- *resp failure
- *core-pulmonale
- *brain abscess
- *amylodiosis.







Comparison of bronchiectasis and chronic obstructive <u>pulmonary disease (COPD)</u>

<u>Bronchiectasis</u>

Etiology Infection or genetic or immune defect

Cigarette smoking

Role of infection Primary

Secondary

Predominant organism in sputum

Haemophilus influenzae, ž

Pseudomonas aeruginosa

Streptococcus pneumoniae,

H. influenzae

Airflow obstruction and \geq

hyperresponsiveness Present

Present

Findings on chest imaging Airway dilation and thickening, ž

mucous plugs

Hyperlucency,

hyperinflation, airway dilation

Quality of sputum Purulent, three-layered

Mucoid, clear