Expiratory central airway collapse

Exaggerated luminal narrowing during expiration due to:

- Excessive dynamic airway collapse of the posterior membrane (EDAC)
- Pathological collapse of the cartilaginous rings
 - o Tracheomalacia (TM): limited to the trachea
 - Tracheobronchomalacia (TBM): collapse extends into one or both mainstem bronchus
- Combined TM or TBM and EDAC

Etiology

- Congenital
 - Polychondritis
 - o Mounier-Kuhn syndrome "idiopathic giant trachea"
- Acquired
 - Chronic inflammatory lung diseases
 - COPD/asthma, ILD, bronchiectasis/chronic bronchitis, recurrent aspiration/GERD, relapsing polychondritis, AAV
 - o Posttraumatic
 - Intubation, tracheostomy, chest trauma
 - o Chronic external compression
 - o Obesity makes the expiratory airways alterations more prominent

Classification based on tracheal narrowing

- Mild: 70 to 80%
- Moderate: 81 to 90%
- Severe: >90%

Clinical presentation

TM-TBM and EDAC have similar clinical presentation, diagnostic evaluation, and share some therapeutic measures.

Mild to moderate are usually asymptomatic and moderate to severe are commonly associated with:

- Dyspnea
- Cough, frequently barking quality
- Inability to clear airway secretions
 - Wheezing
 - Recurrent bronchitis/pneumonia

Diagnosis

- Direct visualization by bronchoscopy
- Dynamic airway CT scan can be used to measure the airway collapsibility index (ACI)
 - Area at the end of inspiration (AEI) minus dynamic area at the end of expiration (DEA) divided by the AEI x100 (AEI-DEA/AEI x100)

Treatment:

- Asymptomatic
 - Management of underlying conditions and observation

- Because it is usually progressive, it is recommended for patient >60 years frequent assessment:
 - PFTs with flow-volume loop yearly
 - Dynamic airway CT chest every two years
- Symptomatic
 - Management of underlying conditions
 - Pursed lip breathing
 - o Promote airway clearance
 - Oscillating positive expiratory pressure (OPEP) q12h (breath through nose and a hold for 2 sec and repeat for up to 10 min)
 - Albuterol followed by saline 3% or 7% or NAC 10 0r 20% by nebulization q12h or daily
 - Pulmonary rehabilitation
 - Baseline assessment with reassessment in 4 to 8 weeks:
 - Clinical parameters (mMRC dyspnea score, cough, and ability to clear secretions
 - Spirometry with flow-volume loop
 - 6 MWT
 - Consider beneficial response if both:
 - Improvement in two out of three of the clinical parameters
 - Improvement in spirometry and or 6MWT
 - If beneficial response continue with medical therapy
 - o If no beneficial response assess suitability for stent trial for 1-2 weeks if:
 - TBM is severe and central (limited to trachea, main bronchus and bronchus intermedius)
 - Persistent or worsening dyspnea
 - Stents can improve dyspnea but usually no other symptoms
 - No large tracheomegaly
 - \circ $\:$ If dyspnea does not improve, TBM is unlikely to be the cause
 - If improvement and patient can tolerate surgery, refer for central airway stabilization surgery
 - Tracheobronchoplasty, open or robotic
 - Less frequently short-segment tracheal resection if suitable
 - If patient is not a surgical candidate
 - NIV (CPAP or BiPAP). Can maintain an open airway and facilitate secretion drainage
 - Bronchoscopy with a PAP facial mask in place
 - $_{\odot}~$ PAP pressures are increased until airway collapse is <70% or a pressure of 16 cm H_2O is reached
 - If bronchoscopic titration is not feasible, start PAP at 8 cm H₂O and titrate up according to symptoms
 - For patients with hypercapnic respiratory failure use BiPAP

- Initially PAP 24 hours per day with gradual transition to intermittent PAP as tolerated (usually while sleeping and daytime as needed along with pursed lip breathing
- Long-term stenting only if NIV fails
- Tracheostomy as a last resort for patients with respiratory failure requiring NIV