Chronic Obstructive Pulmonary Disease (COPD)

These Clinical Practice Guidelines are guidelines only. In no way should these be used as a substitute for clinical or medical judgment. For specialty patient populations such as elderly or post-partum patients, refer to evidenced based practice guidelines to best serve these populations' unique needs.

COPD Quick Facts

- COPD is the third leading cause of death in the U.S.
- COPD accounts for 20% of hospitalized patients over the age of 40.
- 30 day readmission rates are high (20.6%), second only to heart failure
- Frequently identified causes of COPD readmissions:
 - Inability to afford medications
 - Incorrect use of inhalers and nebulizers
 - Lack of transportation to post-discharge appointments

Clinical Concern for COPD Diagnosis

In patients presenting with any of the following symptoms, consider the diagnosis of COPD and obtain spirometry:

- Patients with dyspnea that is persistent, worse with exercise, & progressive over time
- Recurrent wheezing
- Chronic cough: continuous or intermittent, dry or productive
- Recurrent lower respiratory infections
- History of risk factors: tobacco smoke, home or environmental smoke (cooking or heating fuels, occupational exposures)
- Host factors: prematurity, childhood respiratory infections, genetic factors, developmental abnormalities

General questions regarding COPD management or whether a referral to Pulmonary Medicine is needed?

Email <u>COPDhelpline@uhhospitals.org</u>

This account is monitored Monday – Friday by Pulmonary APPs to offer guidance in the management of COPD patients.

It is NOT for URGENT or EMERGENT ISSUES. Our goal is to send a response within 48 to 72 hours.







Capturing the Patient's Perception of Breathlessness and Life Impact of COPD Over Time

Assess at each visit:

- The Modified Medical Research Council Dyspnea Scale (mMRC)
- The COPD Assessment Test (CAT)

The mMRC and CAT assessment tools can be used pre-COPD to gauge symptoms and after a diagnosis of COPD, should be assessed at each visit.

| Modified MR | C Dyspnea Sca | ale | | Table 2.7 |
|--|--|--|---|--|
| PLEASE TICK IN TH | E BOX THAT APPLI | ES TO YOU ONE BO | OX ONLY Grades 0 |) - 4 |
| mMRC Grade 0 | mMRC Grade 1 | mMRC Grade 2 | mMRC Grade 3 | mMRC Grade 4 |
| I only get breathless with strenuous exercise | I get short of breath when hurrying on the level or walking up a slight hill | I walk slower than people of the same age on the level because of breathlessness, or I have to stop for breath when walking on my own pace on the level | I stop for breath after walking about 100 meters or after a few minutes on the level | I am too breathless to leave the house or I am breathless when dressing or undressing |
| | | | | |
| Reference: ATS (1982) Am Rev Respir Dis. Nov;126(5):952-6. | | | | |

Figure from Reference #3 on page 16.

| CAT™ Assessment | | | Figure 2. |
|--|-------------|--|-----------|
| For each item below, place a mark Be sure to only select one response | | cribes you currently. | |
| EXAMPLE: I am very happy | 0 🗶 2 3 4 5 | I am very sad | Score |
| I never cough | 012345 | I cough all the time | |
| I have no phlegm (mucus) in my chest at all | 012345 | My chest is completely full of phlegm (mucus) | |
| My chest does not feel tight at all | 012345 | My chest feels very tight | |
| When I walk up a hill or one flight of stairs I am not breathless | 012345 | When I walk up a hill or one flight of stairs I am very breathless | |
| I am not limited doing any activities at home | 012345 | I am very limited doing activities at home | |
| I am confident leaving my home despite my lung condition | 012345 | I am not at all confident leaving my home because of my lung condition | |
| I sleep soundly | 012345 | I don't sleep soundly because of my lung condition | |
| I have lots of energy | 012345 | I have no energy at all | |
| | | | \square |

Figure from Reference #3 on page 16.



Start with evaluation using the Pulmonary Function Test (PFT) and 6 Minute Walk Test (6MWT). Order "Pre-post Bronchodilator Spirometry".

Pulmonary Function Test: Required for Diagnosis

- Obstruction when the post bronchodilator is FEV1/FVC < LLN or 0.7 (70%)
- FEV1% predicted
 - >80% GOLD 1 (mild)
 - 50 79% GOLD 2 (moderate)
 - 30-49% GOLD 3 (severe)
 - < 30% GOLD 4 (very severe)

6 Minute Walk Test:

- Measures achieved walking distance, dyspnea, and vital signs to assess patient's cardiopulmonary and musculoskeletal response to exercise
- Can be used to evaluate exercise capacity and initiate or titrate oxygen if needed

PFTs can be repeated every 3 years if the patient is stable and not smoking. PFTs can be repeated yearly if needed for therapeutic decision making (medication management, identify or rule out new pulmonary diagnoses) or if there is a suspicion for significant disease progression.

After the Diagnosis

Determine the GRADE (severity) (1-4) and GROUP (staging) (A,B,E) to understand severity and guide medication selection. GOLD ABE assessment requires PFT results, the mMRC and CAT assessments and the number of exacerbations and/or hospitalizations within the last 12 months.



Figure from Reference #3 on page 16.



Approach to the Overall Management of COPD 🛕



Figure from Reference #3 on page 16.

Living Well With COPD Booklet

Order #807987 Ensure patient recieves the booklet as soon as possible





Pharmacological Treatment

Consider ICS use:

- Favors Use: History of ECOPD (COPD with exacerbations) with hospitalizations, ≥ 2 moderate ECOPD/year, blood eosinophils ≥ 300 cells/µL, concomitant asthma
- Against: Repeated pneumonia events, blood eosinophils < 100 cells/µL, history of mycobacterial infection

| ≥ 2 moderate exacerbations or ≥ 1 leading to hospitalization | | LAMA* HCS* if blood eos ≥ 300 |
|---|-----------------------------|----------------------------------|
| 0 or 1 moderate exacerbations (not leading to hospital admission) | GROUP A A bronchodilator | GROUP B LABA + LAMA* |
| mMRC 0-1, CAT < 10 mMRC ≥ 2, CAT ≥ 10 | | |

Inhaler Classes: Initial Treatment

All groups (A, B, E) require a SABA PRN.

Group A

Scheduled LAMA or LABA
 Short acting SABA or SAMA alone

Group B

1. LAMA + LABA (can be combination)

Group E

1. LAMA + LABA 2. LAMA + LABA + ICS

| Drug Names | | |
|------------------|---|--|
| LABA | Long-acting beta agonist | |
| LAMA | Long-acting muscarinic antagonist | |
| SABA and SAMA | Both are the same except "short- acting" | |
| ICS | Inhaled corticosteroid | |



Commonly Used Inhalers in the United States

| Class | Medication in Class Generic (Brand, Inhaler type/name) | Frequency | t _{1/2} | Duration | |
|--|--|--|--|--|--|
| Single Agents | Single Agents | | | | |
| Inhaled Corticosteroids (ICS) | Belomethasone (Qvar RediHaler, MDI) Budesonide (Pulmicort Flexhaler ^e , neb susp ^{ud}) Ciclesonide (Alvesco MDI) Fluticasone (Arnuity Ellipta ^e , Flovent HFA, Diskus ^e) Mometasone (Asmanex HFA) | QD BID BID QD-BID QD-BID | | | |
| Long-Acting Beta Agonists (LABA) | Arformoterol (Brovana neb soln) Formoterol (Foradil aerolizer ^{DC} /Perforomist neb soln ^{ud}) Indacaterol (Arcapta Neohaler ^{DC}) Olodaterol (Striverdi Respimat) Salmeterol (Serevent Diskus ^P) | BID BID - QD BID | 26 h 7h ^{neb} - 7.5 h 5.5 h | ? 12 h - 24 h 12 h | |
| Long-Acting Muscarinic Antagonists (LAMA) | Aclidinium (Tudorza Pressair ^p) Tiotropium (Spiriva Handihaler ^p , Respimat) Umeclidinium (Incruse Ellipta ^p) | BID QD QD | 5-8 h 25-44h 11 h | ? ? ? | |
| Short-Acting Beta Agonists (SABA) | Albuterol (Ventolin HFA, Proventil HFA, ProAir HFA, Digihaler [®] , RespiClick [®] , neb soln) Levalbuterol (Xopenex HFA, neb soln) Pirbuterol (Maxair Autohaler ^{DC}) | q4-6h q4-6h ^{mdi} q6-8h _{neb} - | 3.8-5 h 3.3-4 h - | 4-6h ^{mdi} 3-6h ^{neb} 3-4 h ^{mdi} 5-6h ^{neb} - | |
| Short-Acting Muscarinic Antagonist (SAMA) | Ipratropium (Atrovent HFA) Ipratropium (Atrovent neb soln) | q4-6h ^{mdi} q6-8h ^{neb} | 2 h | 2-4h ^{mdi} 4-5h ^{neb} | |
| Combination Agent | ts | | • | • | |
| ICS + LABA A | Budesonide and Formoterol (Symbicort MDI) Fluticasone and Salmeterol (Advair HFA, Diskus [®] , AirDuo Digihaler [®] , AirDuo RespiClick [®] , Wixela Inhub [®]) Fluticasone and Vilanterol (Breo Ellipta) Mometasone and Formoterol (Dulera MDI) | BID BID | | | |
| | | QD BID | | | |
| | CS + LAMA + ABA Fluticasone, Umeclidinium, Vilanterol (Trelegy Ellipta ^P) Budesonide, Glycopyrrolate, Formoterol (Breztri Aeropshere) | QD | | | |
| LABA | | BID | | | |
| LAMA + LABA | Aclidinium and Formoterol (Duaklir Pressair [®]) Tiotropium and Olodaterol (Stiolto Respimat) Umeclidinium and Vilanterol (Anoro Ellipta [®]) | BID QD QD | | | |
| SAMA + SABA | Ipratropium and Albuterol (Combivent Respimat) Ipratropium and Albuterol (Duoneb neb soln) | q4-6h | | | |





Figure from Reference #9 on page 16.

Choosing the Appropriate Inhalation Device

- Patient dexterity, strength, coordination, and cognition must be evaluated
- Try to minimize the number and types of devices
- Dry powder inhalers (DPI) require patients to make a forceful and deep inhalation
- Metered-dose inhalers (MDI) require coordination between device triggering & inhalation (needs to be slow and deep)
- Adding a spacer (a holding chamber) to an MDI can help with delivery
- Soft mist inhalers (SMI) release the drug as a fine mist more slowly than an MDI to help with a slower inhalation
- Patients who have difficulty with DPI, MDI or SMI inhalers should be considered for nebulizer delivery
- Some patients can be sensitive to DPIs with side effects of sore throat or hoarse voice.

Inhaler Teaching

- Assess for barriers to getting and using inhalers
- <u>COPD Foundation APP</u> has great inhaler and nebulizer educational videos





Address with the Initial and Ongoing Assessment:

Smoking History

- 1. Status: Current, Never, Former + Quit Date
- 2. Pack years:
 - a.1 ppd x 20 yrs = 20 pack years
 - b. 0.5 ppd X 20 yrs = 10 pack years

Smoking Cessation

- 1. Education done documenting "> 5 minutes of smoking cessation counseling"
- 2. Give patient "How to be a Quitter" book
- 3. Referral to Tobacco Cessation Counselor, for questions email <u>Tobacco.Treatment@UHhospitals.org</u>

Immunizations for Adults with COPD

Follow the CDC Adult Immunization Schedule.

Specifically recommended for COPD:

- Influenza
- Pneumonia: Pneumococcal: 2 options CDC recommendation

 1 dose 20 valent (PCV20) OR
 - 1 dose 15 valent (PCV15) followed by 23-valent (PPSV23)
- Tdap (dTaP/dTPa)
- COVID
- Shingles

Lung Cancer Screening

- In order to qualify:
 - Age 50-77
 - 20 pack years +
 - Smoked within the last 15 years
 - Shared decision making discussion completed & documented
 - No previous CT chest in the last year (includes Ca scoring)
- Ordered as CT chest low dose for lung screening w/o contrast
- If original screening CT is abnormal and requires sooner follow up based on Lung RADS score- need to order "CT Chest Lung CA Screen Follow Up LR3LR4 Diagnostic"
- Only cigarette smokers qualify for this program. It does not take into account other types of tobacco/ smoking products cigars, vape, or marijuana.
- Lung cancer screening navigators <u>UHLungCancerScreeningProgram@UHhospitals.org</u>

Oxygen: Current Requirements & Oxygen Evaluation

Goal Pulse Ox (88% - 92%)

Long term oxygen therapy indicated for:

- 1. PaO2 ≤ 55 mmHg or SaO2 ≤ 88% with or without hypercapnia, confirmed twice over a 3 week period OR PaO2 55-60 mmHg or SaO2 88% if there is evidence of pulmonary hypertension, peripheral edema suggesting CHF or polycythemia (hematocrit > 55%)
- 2. Re-evaluate at 60-90 days with ABG or oxygen saturation measurements: room air AND on prescribed O2 to assess for continued need and flow requirements
- 3.O2 desaturation study and provider visit within 30 days required for oxygen initial certification and annually for recertification





Exercise and Self-Management

Current level of physical activity? Understanding of breathlessness? Utilization of techniques to manage anxiety / SOB?

For self-management:

- Avoid infection
- Maintain good nutrition
- Aerobic exercise can refer to Pulmonary Rehabilitation, if they qualify

Pulmonary Rehabilitation

- Groups B or E (or restriction on PFTs or low DLCO)
- Patient is allotted 72 sessions (lifetime)

ICD-10 Codes That Support Pulmonary Rehabilitation

- D86.9 Sarcoidosis
- E84.0 Cystic Fibrosis with pulmonary manifestations
- I27.0 Primary pulmonary hypertension
- J41.1 Simple chronic bronchitis
- J44.9 Obstructive chronic bronchitis without acute exacerbation
- J41.1 Other chronic bronchitis
- J43.9 Other Emphysema
- J44.9 Chronic Obstructive Asthma unspecified
- J45.991 Cough-variant asthma
- J47.9 Bronchiectasis without acute exacerbation
- J44.9 Chronic airway obstruction not elsewhere classified (COPD)
- J60 Coal workers' pneumoconiosis
- J61 Asbestosis
- J62.8 Pneumoconiosis due to other silica or silicates
- J63.6 Pneumoconiosis due to other inorganic dust
- J66.8 Pneumonopathy due to inhalation of other dust
- J64 Pneumoconiosis unspecified
- J68.4 Chronic respiratory conditions due to fumes and vapors
- J68.9 Unspecified respiratory conditions due to fumes and vapors
- J70.1 Chronic and other pulmonary manifestations due to radiation
- J84.10 Postinflammatory pulmonary fibrosis
- J84.01 Pulmonary alveolar proteinosis
- J84.02 Pulmonary alveolar microlithiasis
- J84.09 Other specified alveolar and parietoalveolar pneumonopathies
- J98.4 Other diseases of lung not elsewhere classified

Additional tips/ helpful hints:

- In order for COPD to qualify patient needs to have a moderate obstruction (FEV1 50-80%).
- Non-COPD dx require FVC < 65% or FEV1 < 65% or DLCO < 65%
- Pulmonary hypertension and long haul COVID qualifies
- Each pulmonary rehab referral is for 36 sessions. If they want to utilize the additional 36 sessions (72 lifetime sessions) they need to be referred again.
- Pulmonary rehabilitation can ONLY be ordered by a physician (APP [NP/PA] must put supervised by a collaborating physician in the order)



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Addressing Comorbidities and When to Refer to Pulmonary/Sleep Medicine

Comorbidities

1. Identification & optimal management including referrals, as needed, is the goal

- 2. Common Comorbidities:
 - Hypertension, Obstructive Sleep Apnea, Gastroesophageal Reflux
 - Cardiovascular Disease, Heart Failure, Arrhythmias
 - Pulmonary Embolism, Pulmonary Hypertension
 - Osteoporosis, Anxiety, Depression

Obstructive Sleep Apnea (OSA)

Patients with COPD and OSA have a worse prognosis and are more likely to develop daytime pulmonary hypertension than either condition alone. In patients with both COPD and OSA, there are clear indications for the use of CPAP: reduced all-cause hospitalizations, ED visits, moderate and severe exacerbations, and associated healthcare costs.

To screen for OSA using the STOP BANG instrument, see the below diagram as well as the UHQCN Sleep CPG.

| STOP | BANG | |
|---|--|--|
| Do you SNORE loudly? Do often feel TIRED? Has anyone OBSERVED you stop breathing? Do you have/are you treated for high blood PRESSURE? | BMI >35 AGE>50 NECK circumference (>17 in men, >16 in women) Gender= male | STOP BANG Scoring 0-2 Low 3-4 Intermediate 5-8 High |

| Low Pretest Probability | Moderate Pretest Probability | High Pretest Probability |
|---|--|--------------------------|
| • STOP BANG < 3 | STOP BANG 3-4 Uncontrolled blood pressure ↑ test need | • STOP BANG 5-8 |
| Try conservative strategies for 3 months; then reassess need to test. | Appropriate to test; Use clinical judgement. | High priority to test |

Pulmonary Medicine Referral

- H Hospitalizations ≥ 1 exacerbation per year
- E Exacerbations ≥2 requiring ED visits/steroids
- L Low lung function. FEV1 < 50% (GOLD 3 or 4)
- P Problem sleeping or symptoms of sleep disorder breathing
- O Oxygen needs
- **U** Uncertain diagnosis
- T Therapy options (need to up-titrate)



Exacerbations in COPD

COPD Exacerbation

Defined as an event characterized by dyspnea and/or cough and sputum that worsens in < 14 days

Cardinal Symptoms: dyspnea, sputum volume, sputum purulence*

- 1. Mild worsening of 1 of the cardinal symptoms
 - RR < 24, HR < 95, dyspnea < 5 on VAS
 - Resting sat \geq 92% on room air/usual O2 (or change \leq 3% from baseline)
- 2. Moderate worsening of 2 of the 3 cardinal symptoms
 - RR ≥ 24, HR ≥ 95
 - Resting sat < 92% on RA/usual O2 (or change > 3% from baseline)
 - ABG may show hypoxemia &/or hypercapnia but NOT acidosis
- 3. Severe worsening of all 3 cardinal symptoms
 - Parameters same as moderate
 - ABG shows hypercapnia & acidosis

*Antibiotics should only be given if sputum purulence is one of the presenting symptoms

Make sure the patient has a rescue inhaler: SABA or SAMA/SABA.

| Exacerbation: Outpatient Treatment Plan | | |
|---|--|--|
| Exacerbation Level | In all cases, re-evaluate in 48 hours for persistent or worsening symptoms and the need to escalate treatment. | |
| Mild | Increase use of PRN albuterol/nebulizers to every 4 hours Continue long acting (maintenance) inhalers | |
| Moderate | Prednisone 40mg daily x 5 days Antibiotics (if + sputum purulence) see the next page Increase use of PRN albuterol/nebulizers to every 4 hours Continue long acting (maintenance) inhalers | |
| Severe | Prednisone 40mg daily x 5 days Antibiotics (see below) Evaluate for referral to ED Increase use of PRN albuterol/nebulizers to every 4 hours Continue long acting (maintenance) inhalers | |





A Guide to Antibiotics

| Clinical Indicators (Exacerbation of COPD: ECOPD) | Antibiotic Options Recommended duration of treatment for outpatient ECOPD is ≤ 5 days |
|---|--|
| FEV ¹ > 50% AND ≤ 2 ECOPD / year | Macrolide OR 2nd/3rd generation cephalosporin OR Trimethoprim / Sulfamethoxazole |
| FEV ¹ < 50% >2 ECOPD / year ECOPD with hospitalization in the last 12 months Home O ² Comorbidities: heart failure, ischemic heart disease Chronic oral steroid use Antibiotic use in the last 3 months | Fluoroquinolone OR Amoxicillin / Clavulanate |
| FEV ¹ < 30% Bronchiectasis Known Pseudomonas Chronic oral steroids Multiple risk factors | • Fluoroquinolone |

| Commons Antibiotics This is not an exhaustive and please contact a Pharmacist with questions. | | |
|---|--|--|
| Macrolide | AzithromycinErythromycinClarithromycin | |
| 2nd Generation Cephalosporin | CefuroximeCefonicidCeforanide | |
| 3rd Generation Cephalosporin | CefiximeCeftibutenCefdinir | |
| Fluoroquinolone | CiprofloxacinLevofloxacinGemifloxacin | |





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Possible Hospitalization of Emergency Department Treatment

SEND to the Emergency Department...

If any of the following:

- Marked increased intensity of symptoms
- New or worsening peripheral edema
- Worsening of hypoxemia from usual (if known)
- SpO2 <92% if not on home oxygen
- Shortness of breath that is worsening and/or at rest and is not relieved or controlled with the use of rescue inhaler/nebulizer treatment
- High fever
- Altered mental state (confusion, slurred speech, drowsiness)
- Chest pain
- Worsening of co-morbidities (e.g. heart failure, ischemic heart disease, diabetes)
- Inability to perform daily activities and/or manage safely at home
- Increased anxiety (feeling scared/afraid)





Advanced COPD Recommendations and Interventions

1. Additional COPD Medications

- Roflumilast (Daliresp) if FEV1 < 50% predicted and chronic broncititis
- Macrolide (Azithromycin)
 - Monitor QTc
 - Avoid in patients with cardiac disease/arrhythmias
 - Can cause hearing loss
- Withdraw ICS if recurrent pneumonia episodes

2. Advance Directives & Palliative Medicine Referral

- Advance directives include living will and durable power of attorney for healthcare
- End-of-life care should be discussed with patients and families to understand their thoughts and beliefs regarding what is an acceptable quality of life, desired level of resuscitation, place of death preferences and creating advanced directives.
- Palliative's role is to improve symptom management and assist the patient to achieve the best possible quality of life, including facilitating goals of care conversations and decisions.
- Even with optimal therapy many patients experience breathlessness, reduced exercise capacity, fatigue, anxiety, panic attacks and depression.
- Refer COPD patients to palliative medicine when they are O2-dependent PLUS one or more of the following:
 - Age > 85
 - Nutritional decline (>10% weight loss over past year)
 - 2+ hospitalizations or ICU admission in the past 6 months
 - $\circ~$ Additional organ system failure dementia, CHF with severely reduced EF, CKD IV or V, or cirrhosis with MELD > 15

3. Non-Invasive Ventilation (NIV)

NIV has several variations. NIV can provide single level pressure support (EPAP) or bi-level support (IPAP and EPAP). That support can also provide a back-up respiratory rate or none at all. Patients with severe COPD (Gold 3-4, FEV1 < 50% predicted), daytime fatigue, dyspnea, suspicion of hypercapnia, and a history of hospitalization for acute respiratory failure should be evaluated for the use of NIV.

Nocturnal noninvasive, ventilation, (CPAP, BIPAP)

The first step is to get an ABG and confirm chronic hypercapnia PaCO2 > 52 mmHg.

Patients with severe COPD and daytime fatigue without chronic hypercapnia should be referred to sleep medicine for a polysomnogram (sleep study) as there is a potential for OSA/COPD overlap that may require CPAP support.

Patients with severe COPD with confirmed hypercapnia:

- 1. Obtain an overnight oximetry study, looking for evidence of hypoxemia.
- 2. The study is completed either on the patient's prescribed home O2 or 2 L per minute; whichever is greater.
- 3. If the patient (while on the oxygen) has a saturation < 88% for 5 minutes; (not continuous), the patient has demonstrated severe COPD with hypoxemia and is qualified to receive NIV without a backup rate.
- 4. Per the CMS guidelines this is known as a Respiratory Assist Device (RAD) without a backup rate. The goal is to use bi-level support with the IPAP titrated to reach > 18 cm H2O and an exhaled tidal volume of 8mL/kg IBW.





Proportional Open Ventilation (POV)

POV is noninvasive ventilation that is portable, multi modal, and can work with or without supplemental oxygen. It is designed for patients with chronic hypoxic respiratory failure who have a significant amount of symptoms throughout the day, negatively impacting their ability to function. POV delivers breaths in various modes and according to the type of activity, supporting overall ventilation, reducing work of breathing and improving quality of life.

The best example of this type of ventilation is the life 2000 ventilation system (manufactured by Hillrom). There is a stationary component in the home as well as a small, portable volume ventilator worn around the waist with a six hour battery that weighs approximately 1 pound. Delivery of breaths is accomplished through a nasal pillow. Candidates for Life 2000 have chronic respiratory failure as in COPD, restrictive lung disease, neuromuscular disorders, and lung transplant (pre and post). When using this device for COPD, the ultimate goal is reduce exacerbations and improve quality of life.

4. Alpha1-Antitrypsin Deficiency (AATD)

- 1. WHO recommends all COPD patients be screened for AATD
- 2. Additionally consider screening patients with liver disease of unknown cause, unexplained bronchiectasis, ANCA positive vasculitis or first degree relatives of an affected individual
- 3. Obtain AAT level AND targeted genotype (see algorithm in UpToDate)
 - i. AAT blood level low if < 20 micromol/L or < 100 mg/dL
 - ii. Look for presence of deficient variant (F, I, S, Z)
 - iii. If no gene sequence of SERPINA1 looking for null or rare variants

6. Lung Volume Reduction Surgery/Endobronchial Valve (Zephyr Valve)

COPD patients with emphysema and significant hyperinflation refractory to optimal medical care

Inclusion Criteria:

- Diagnosis of emphysema confirmed by CT
- BMI <35
- Stable with < 20mg prednisone (or equivalent) daily
- RV > 175% predicted (>200% if homogenous)
- FEV1 15-45% predicted
- TLC >100% predicted
- Not actively smoking (for at least 4 months)
- 6 minute walk distance 100-500m (150-500 if homogenous)

Exclusion Criteria:

- Currently smoking
- Prior lung transplant, lung volume reduction surgery, or lobectomy
- Prior surgery requiring median sternotomy (CABG etc.)
- CHF EF <45%, unstable cardiac arrhythmia, MI or CVA
- Allergies to nitinol, nickel, titanium, or silicone
- Large bulla >30% of either lung
- Inability to complete preoperative/ postoperative pulmonary diagnostic and therapeutic program required for procedure
- Contraindications for bronchoscopy patient characteristics that may carry a high risk of postoperative morbidity and/or mortality
- Severe hypercapnia (PaCO2 >50 mmHg on RA) and or severe hypoxemia (PaO2 <45 mmHg on RA)
- Uncontrolled pulmonary hypertension (sPAP > 45mmHg)

For any inquiries or referrals for endobronchial valves please contact Toby Schweinfurth RN, Bronchoscopy Coordinator - <u>Toby Schweinfurth@UHhospitals.org</u>

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6.Clinical Trials for COPD

Pulmonology does participate in clinical trials for COPD. In order the information to be accurate, we ask that you search using the link below.

https://www.uhhospitals.org/uh-research/find-clinical-trials-and-studies

7. Lung Transplant Evaluation

- Transplant referral is managed by the patient's Pulmonologist
- Transplant for COPD is primarily for better quality of life and is an option for patients with progressive disease despite maximal therapy that includes medication, pulmonary rehabilitation and supplemental oxygen
- Patients who are appropriate for referral should meet all of the following criteria
 - A BODE index score ≥7. The BODE score is an index of disease severity using Body mass index, airflow Obstruction, Dyspnea & Exercise capacity.
 - Postbronchodilator forced expiratory volume in one second (FEV1) <15 percent of predicted
 - Evidence of hypercapnia defined as carbon dioxide tension (PaCO2) >50 mmHg (6.6 kPa)
 - Not candidate for lung volume reduction surgery
- COPD patients with moderate to severe Pulmonary Artery Hypertension should be considered for lung transplant sooner rather than later



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