



An Independent Licensee of the Blue Cross Blue Shield Association

PHARMACY COVERAGE GUIDELINES  
SECTION: DRUGS

ORIGINAL EFFECTIVE DATE: 2/21/2019  
LAST REVIEW DATE: 2/18/2021  
LAST CRITERIA REVISION DATE:  
ARCHIVE DATE:

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## YUPELRI™ (revefenacin) oral inhalation solution

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Coverage for services, procedures, medical devices and drugs are dependent upon benefit eligibility as outlined in the member's specific benefit plan. This Pharmacy Coverage Guideline must be read in its entirety to determine coverage eligibility, if any.

This Pharmacy Coverage Guideline provides information related to coverage determinations only and does not imply that a service or treatment is clinically appropriate or inappropriate. The provider and the member are responsible for all decisions regarding the appropriateness of care. Providers should provide BCBSAZ complete medical rationale when requesting any exceptions to these guidelines.

The section identified as "Description" defines or describes a service, procedure, medical device or drug and is in no way intended as a statement of medical necessity and/or coverage.

The section identified as "Criteria" defines criteria to determine whether a service, procedure, medical device or drug is considered medically necessary or experimental or investigational.

State or federal mandates, e.g., FEP program, may dictate that any drug, device or biological product approved by the U.S. Food and Drug Administration (FDA) may not be considered experimental or investigational and thus the drug, device or biological product may be assessed only on the basis of medical necessity.

Pharmacy Coverage Guidelines are subject to change as new information becomes available.

For purposes of this Pharmacy Coverage Guideline, the terms "experimental" and "investigational" are considered to be interchangeable.

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This Pharmacy Coverage Guideline does not apply to FEP or other states' Blues Plans.

Information about medications that require precertification is available at [www.azblue.com/pharmacy](http://www.azblue.com/pharmacy).

Some large (100+) benefit plan groups may customize certain benefits, including adding or deleting precertification requirements.

All applicable benefit plan provisions apply, e.g., waiting periods, limitations, exclusions, waivers and benefit maximums.

Precertification for medication(s) or product(s) indicated in this guideline requires completion of the [request form](#) in its entirety with the chart notes as documentation. **All requested data must be provided.** Once completed the form must be signed by the prescribing provider and faxed back to BCBSAZ Pharmacy Management at (602) 864-3126 or emailed to [Pharmacyprecert@azblue.com](mailto:Pharmacyprecert@azblue.com). **Incomplete forms or forms without the chart notes will be returned.**



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### Criteria:

- **Criteria for initial therapy:** Yupelri (revefenacin) is considered *medically necessary* and will be approved when **ALL** of the following criteria are met:
1. Individual is 18 years of age or older
  2. A confirmed diagnosis of chronic obstructive pulmonary disease (COPD)
  3. Individual has failure, contraindication or intolerance to **two** trials of the following inhaled anticholinergic/anti-muscarinic with an inhaled long-acting beta-agonist with or without an inhaled corticosteroid:
    - a. Incruse ellipta (umeclidinium)
    - b. Seebri neohaler (glycopyrrlate)
    - c. Spiriva (tiotropium)
    - d. Tudorza (aclidinium)
  4. Individual has failure, contraindication or intolerance to ipratropium bromide solution for nebulization
  5. Individual does not have hepatic impairment
  6. Individual is a non-smoker or is quitting through use of behavior modification and/or medications aimed at smoking cessation
  7. There are **NO** contraindications
    - a. Contraindications include:
      - i. Hypersensitivity to revefenacin or any component of the product

**Initial approval duration:** 6 months

- **Criteria for continuation of coverage (renewal request):** Yupelri (revefenacin) is considered *medically necessary* and will be approved when **ALL** of the following criteria are met:
1. Individual's condition has responded
    - a. Response is defined as BOTH of the following:
      - i. Improved FEV1 over baseline
      - ii. Reduced number and frequency of exacerbations
  2. Individual has been adherent with the medication
  3. Individual is a non-smoker or is quitting through use of behavior modification and/or medications aimed at smoking cessation
  4. Individual does not have hepatic impairment



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5. Individual has not developed any contraindications or other significant level 4 adverse drug effects that may exclude continued use
  - a. Contraindications as listed in the criteria for initial therapy section
  - b. Significant adverse effect such as:
    - i. Paradoxical bronchospasm
6. There are no significant interacting drugs

**Renewal duration:** 12 months

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### **Description:**

Yupelri (revefenacin) inhalation solution is indicated for the maintenance treatment of patients with chronic obstructive pulmonary disease (COPD).

Yupelri (revefenacin) is a long-acting muscarinic antagonist (LAMA). It has similar affinity to the subtypes of muscarinic receptors M1-M5. In the airways, it exhibits pharmacological effects through inhibition of M3 receptor at the smooth muscle leading to bronchodilation. The safety and efficacy of Yupelri (revefenacin) have been established in clinical trials when administered using the PARI LC® Sprint nebulizer with a mouthpiece and the PARI Trek® S compressor. The safety and efficacy delivered from non-compressor based nebulizer systems have not been established.

Characteristics COPD includes small airways disease (obstructive bronchiolitis) and parenchymal destruction (emphysema). The presence of chronic inflammation causes structural changes and narrowing of the small airways.

No one COPD product adds superior clinical value over alternatives within any pharmacologic class. Guidelines recommend COPD medications by class, not by specific medication. A step-wise approach is used to minimize symptoms and reduce frequency and severity of exacerbations. COPD evidence-based clinical practice guidelines recommend combining medications from various pharmacologic classes for long-term management of COPD in a step-wise fashion as symptoms progress. As of yet, no medication modifies long-term decline in lung function.

Initial management of COPD patients includes either an inhaled long-acting beta agonist (LABA) or a LAMA, both agents relax bronchial smooth muscle. An inhaled corticosteroid (ICS) is used for those patients who are at high risk for exacerbations. Other COPD medications include inhaled short-acting bronchodilators (beta-agonists and antimuscarinic agents), methylxanthines, oral corticosteroids, and phosphodiesterase-4 (PDE-4) inhibitors.

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### Definitions:

#### Global Initiative for Chronic Obstructive Lung Disease (GOLD) assessment:

| GOLD: severity of airflow limitation (based on postbronchodilator FEV1)  |                    |                    |
|--|--------------------|--------------------|
| Stage  | Severity           | FEV1 (%predicted)  |
| <b>In patients with FEV1 / FVC &lt; 0.7</b>  |                    |                    |
| GOLD 1   | Mild               | ≥ 80               |
| GOLD 2   | Moderate           | 50-79              |
| GOLD 3   | Severe             | 30-49              |
| GOLD 4   | Very severe        | < 30               |
| <b>GOLD: Assessment of symptoms and risk for exacerbations</b>   |                    |                    |
| Exacerbations/Hospitalizations   | Symptom assessment |                    |
|  | mMRC 0-1; CAT < 10 | mMRC ≥ 2; CAT ≥ 10 |
| 0-1 exacerbations without hospitalization  | A                  | B                  |
| ≥ 2 exacerbations or ≥ 1 hospitalization   | C                  | D                  |
| A: Low risk, less symptoms<br>B: Low risk, more symptoms<br>C: High risk, less symptoms<br>D: High risk, more symptoms<br><br>CAT: COPD Assessment Test<br>mMRC: modified Medical research Council dyspnea scale |                    |                    |

#### Modified Medical Research Council Dyspnea Scale:

| Grade | Description of breathlessness   |
|-------|---|
| 0     | I only get breathless with strenuous exercise   |
| 1     | I get short of breath when hurrying on level ground or walking up a slight hill   |
| 2     | On level ground, I walk slower than people of the same age because of breathlessness or have to stop for breath when walking at my own pace |
| 3     | I stop for breath after walking about 100 yards or after a few minutes on level ground  |
| 4     | I am too breathless to leave the house or I am breathless when dressing   |

#### COPD Assessment Test:

|   | Circle the number that best describes you |  |
|---|---|--|
| I never cough   | 1 2 3 4 5                                 | I cough all the time   |
| I have no phlegm in my chest at all                               | 1 2 3 4 5                                 | My chest is completely full of phlegm                                  |
| My chest does not feel tight at all                               | 1 2 3 4 5                                 | My chest feels very tight  |
| When I walk up a hill or one flight of stairs I am not breathless | 1 2 3 4 5                                 | When I walk up a hill or one flight of stairs I am very breathless     |
| I am not limited doing any activities at home                     | 1 2 3 4 5                                 | I am very limited doing activities at home                             |
| I am confident leaving my home despite my lung condition          | 1 2 3 4 5                                 | I am not at all confident leaving my home because of my lung condition |
| I sleep soundly   | 1 2 3 4 5                                 | I don't sleep soundly because of my lung condition                     |
| I have lots of energy   | 1 2 3 4 5                                 | I have no energy at all  |

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### Management of Stable COPD based on GOLD ABCD assessment of symptoms and risk of exacerbation:

| Category | Symptoms   | Risk   | Suggested treatment   |
|----------|--|--|---|
| A        | Less symptomatic:<br>Mild or infrequent symptoms (breathless with strenuous exercise or when hurrying on level ground or walking up a slight hill) or CAT <10  | Low:<br>0 or 1 exacerbations in the past year without associated hospitalization     | <b>Recommendation:</b><br>Short-acting bronchodilator or combination of short-acting beta-agonist and anticholinergic (antimuscarinic), as needed.<br><b>Alternative:</b><br>Long-acting bronchodilator if beneficial.  |
| B        | More symptomatic:<br>Moderate to severe symptoms (patient has to walk more slowly than others of same age due to breathlessness, has to stop to catch breath when walking on level ground at own pace, or has more severe breathlessness) or CAT ≥10         | Low:<br>0 or 1 exacerbations in the past year without associated hospitalization     | <b>First choice:</b><br>Regular treatment with a long-acting bronchodilator, either LAMA or LABA, based on symptom relief. Short-acting bronchodilator available for symptom control as needed.<br><b>For persistent symptoms:</b><br>Regular treatment with a combination of LAMA and LABA.  |
| C        | Less symptomatic:<br>Mild or infrequent symptoms (breathless with strenuous exercise or when hurrying on level ground or walking up a slight hill) or CAT <10 <sup>Δ</sup>   | High risk:<br>≥ 2 exacerbations per year with one or more leading to hospitalization | <b>First choice:</b><br>Regular treatment with a LAMA; SABA available for symptom control as needed.<br><b>For further exacerbations:</b><br>Regular treatment with a LAMA plus LABA or (less preferred) LABA plus ICS  |
| D        | More symptomatic:<br>Moderate to severe symptoms (patient has to walk slower than others of same age due to breathlessness, has to stop to catch breath when walking on level ground at own pace, or has more severe breathlessness) <sup>¶</sup> or CAT ≥10 | High risk:<br>≥ 2 exacerbations per year with one or more leading to hospitalization | <b>First choice:</b><br>Regular treatment with combination LABA plus LAMA. LABA plus ICS may be preferred, if features of asthma/COPD overlap. SABA available for symptom control as needed. LAMA alone, if LABA contraindicated.<br><b>For further exacerbations:</b><br>Regular treatment with combination of LAMA plus LABA plus ICS or (less preferred in absence of asthma overlap) switch to LABA plus ICS.<br>If exacerbations continue despite triple therapy, additional options for selected patients include roflumilast (if chronic bronchitis and FEV <sub>1</sub> <50% predicted), theophylline, chronic therapy with a macrolide, and stopping ICS |



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### Resources:

Yupelri (revefenacin) product information, revised by Mylan Specialty, L.P. 05-2019, at DailyMed <http://dailymed.nlm.nih.gov> accessed January 25, 2021.

Ferguson GT, Make B. Stable COPD: initial pharmacologic management. In: UpToDate, Stoller JK, Hollingsworth H (Eds), UpToDate, Waltham MA.: UpToDate Inc. <http://uptodate.com>. Accessed on January 25, 2021.

Stoller JK. COPD exacerbations: Management. In: UpToDate, Barnes PJ, Hollingsworth H (Ed), UpToDate, Waltham MA.: UpToDate Inc. <http://uptodate.com>. Accessed on January 25, 2021.

Ferguson GT, Make B. Management of refractory chronic obstructive pulmonary disease. In: UpToDate, Stoller JK, Hollingsworth H (Eds), UpToDate, Waltham MA.: UpToDate Inc. <http://uptodate.com>. Accessed on January 25, 2021.

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