

## Clinical Policy: Inhaled Agents for Asthma and COPD

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[Revision Log](#)

See [Important Reminder](#) at the end of this policy for important regulatory and legal information.

### Description

The following are inhaled agents for asthma and/or chronic obstructive pulmonary disease (COPD) requiring prior authorization:

- Short acting beta-2 agonist (SABA): albuterol (ProAir<sup>®</sup> Digihaler<sup>®</sup>)
- Inhaled corticosteroid (ICS): budesonide (Pulmicort Respules<sup>®</sup>)\*, ciclesonide (Alvesco<sup>®</sup>), fluticasone (ArmonAir<sup>®</sup> Digihaler<sup>™</sup>), mometasone (Asmanex<sup>®</sup> HFA, Asmanex<sup>®</sup> Twisthaler<sup>®</sup>, Flovent<sup>®</sup> HFA, Flovent<sup>®</sup> Diskus<sup>®</sup>)
- Long acting beta-2 agonist (LABA): arformoterol (Brovana<sup>®</sup>), formoterol (Perforomist)
- Long acting muscarinic antagonist (LAMA): aclidinium bromide (Tudorza<sup>®</sup> Pressair<sup>®</sup>), glycopyrrolate (Seebri<sup>™</sup> Neohaler<sup>®</sup>, Lonhala<sup>®</sup> Magnair<sup>®</sup>), revefenacin (Yupelri<sup>®</sup>)
- Combination ICS/LABA: budesonide/formoterol (Symbicort<sup>®</sup>\*, Symbicort Aerosphere<sup>®</sup>), fluticasone/salmeterol (Advair Diskus<sup>®</sup>\*, Advair HFA<sup>®</sup>\*, AirDuo<sup>®</sup> Digihaler<sup>™</sup>, AirDuo<sup>®</sup> RespiClick<sup>®</sup>), mometasone/formoterol (Dulera<sup>®</sup>)
- Combination LABA/LAMA: aclidinium/formoterol (Duaklir<sup>®</sup> Pressair<sup>®</sup>), glycopyrrolate/formoterol (Bevespi Aerosphere<sup>™</sup>), indacaterol/glycopyrrolate (Utibron<sup>™</sup> Neohaler<sup>®</sup>)
- Phosphodiesterase 3 (PDE3) inhibitor and phosphodiesterase 4 (PDE4) inhibitor: ensifentrine (Ohtuvayre<sup>™</sup>)

\*Generic agents do not require prior authorization.

### FDA Approved Indication(s)

ProAir Digihaler is indicated for the treatment or prevention of bronchospasm in adults, adolescents, and children 4 years of age and older with reversible obstructive airway disease. ProAir Digihaler is also indicated for the prevention of exercise-induced bronchospasm (EIB) in patients 4 years of age and older.

The other inhaled agents are indicated as follows:

Drug Name	Asthma	COPD
ICS		
Alvesco	X (Age ≥ 12 years)	
ArmonAir Digihaler	X (Age ≥ 4 years)	
Asmanex HFA	X (Age ≥ 5 years)	
Asmanex Twisthaler	X (Age ≥ 4 years)	
Pulmicort Respules	X (Age 1-8 years)	
Flovent Diskus, Flovent HFA	X (Age ≥ 4 years)	

Drug Name	Asthma	COPD
<b>LABA</b>		
Brovana		X
Perforomist		X
<b>LAMA</b>		
Lonhala Magnair		X
Seebri Neohaler		X
Tudorza Pressair		X
Yupelri		X
<b>ICS/LABA</b>		
Advair Diskus	X (Age ≥ 4 years)	X
Advair HFA	X (Age ≥ 12 years)	
AirDuo Digihaler	X (Age ≥ 12 years)	
AirDuo RespiClick	X (Age ≥ 12 years)	
Dulera	X (Age ≥ 5 years)	
Symbicort	X (Age ≥ 6 years)	X
Symbicort Aerosphere		X
<b>LABA/LAMA</b>		
Bevespi Aerosphere		X
Duaklir Pressair		X
Utibron Neohaler		X
<b>PDE3/PDE4 Inhibitor</b>		
Ohtuvayre		X

**Policy/Criteria**

*Provider must submit documentation (such as office chart notes, lab results or other clinical information) supporting that member has met all approval criteria.*

It is the policy of health plans affiliated with Centene Corporation<sup>®</sup> that inhaled agents for asthma and COPD are **medically necessary** when the following criteria are met:

**I. Initial Approval Criteria**

**A. Inhaled Agents for Asthma or Chronic Obstructive Pulmonary Disease (must meet all):**

1. Diagnosis of asthma or COPD as FDA-approved for the requested agent (*see FDA Approved Indications section*);
2. Age is one of the following (a or b):
  - a. Asthma: Appropriate per the prescribing information for the requested agent (*see FDA Approved Indications section*);
  - b. COPD: ≥ 18 years;
3. Failure of the following formulary agent(s) at up to maximally indicated doses, unless clinically significant adverse effects are experienced or all are contraindicated:

Requested Agent	Required Step Through Agent(s)
ProAir Digihaler	Two generic albuterol sulfate HFA products, each from a different manufacturer

Requested Agent	Required Step Through Agent(s)
Pulmicort Respules	Age is between 1 to 8 years or documentation supports inability to use inhaler devices <i>AND</i> if request is for brand Pulmicort Respules, medical justification supports inability to use generic Pulmicort Respules (e.g., contraindications to excipients)
Flovent Diskus	Fluticasone propionate diskus (Flovent Diskus authorized generic)
Flovent HFA	Fluticasone propionate HFA (Flovent HFA authorized generic)
<u>All other ICS</u> : Alvesco, ArmonAir Digihaler, Asmanex HFA, Asmanex Twisthaler	Qvar <sup>®</sup> RediHaler <sup>™</sup> <i>AND</i> Pulmicort Flexhaler <sup>™</sup> <i>AND</i> Arnuity <sup>®</sup> Ellipta <sup>®</sup> <i>AND</i> fluticasone propionate HFA (Flovent HFA authorized generic) <i>AND</i> fluticasone propionate diskus (Flovent Diskus authorized generic)
<u>LABA</u> : Brovana, Perforomist	Generic (i.e., formoterol for Perforomist requests, arformoterol for Brovana requests) <i>AND</i> Arcapta <sup>®</sup> Neohaler <sup>®</sup> <i>AND</i> Serevent <sup>®</sup> Diskus <sup>®</sup> <i>AND</i> Striverdi <sup>®</sup> Respimat <sup>®</sup> , unless request is for a nebulized LABA and documentation supports inability to use inhaler devices
<u>LAMA</u> : Lonhala Magnair, Seebri Neohaler, Tudorza Pressair, Yupelri	Incruse <sup>®</sup> Ellipta <sup>®</sup> <i>AND</i> Spiriva <sup>®</sup> Handihaler <sup>®</sup> / Respimat <sup>®</sup> , unless request is for a nebulized LAMA and documentation supports inability to use inhaler devices
Brand Advair Diskus	Medical justification supports inability to use generic fluticasone/salmeterol products (generic Advair Diskus, Wixela <sup>™</sup> Inhub <sup>™</sup> ) (e.g., contraindications to excipients)
Brand Advair HFA	Medical justification supports inability to use fluticasone-salmeterol HFA (Advair HFA authorized generic) (e.g., contraindications to excipients)
Brand Symbicort, Symbicort Aerosphere	Medical justification supports inability to use generic Symbicort (e.g., contraindications to excipients)
<u>All other ICS/LABA</u> : AirDuo Digihaler, AirDuo RespiClick	fluticasone-salmeterol HFA (Advair HFA authorized generic) <i>AND</i> budesonide/formoterol (Symbicort authorized generic) <i>AND</i> fluticasone/salmeterol (generic Advair Diskus or Wixela Inhub) <i>AND</i> Breo Ellipta <sup>®</sup> (brand Breo Ellipta or [fluticasone furoate-vilanterol] Breo Ellipta authorized generic) <i>AND</i> Dulera
<u>LABA/LAMA</u> : Bevespi Aerosphere, Duaklir Pressair, Utibron Neohaler	Anoro <sup>®</sup> Ellipta <sup>®</sup> <i>AND</i> Stiolto <sup>®</sup> Respimat <sup>®</sup>
Ohtuvayre	<ul style="list-style-type: none"> <li>Anoro Ellipta <i>OR</i> Stiolto Respimat <i>OR</i> one LABA (e.g., Serevent Diskus) in combination with one LAMA (e.g., Incruse Ellipta)</li> </ul>

Requested Agent	Required Step Through Agent(s)
	<p><i>AND</i></p> <ul style="list-style-type: none"> <li>• For members with blood eosinophil count <math>\geq 100</math> cells/mcL: Breztri Aerosphere™ <i>OR</i> Trelegy™ Ellipta®</li> </ul> <p><i>Note: Prior failure of triple therapy (ICS/LABA/LAMA) satisfies the requirement for failure of dual therapy (LABA/LAMA).</i></p>

4. For requests for an agent with a digital component (e.g., Digihaler products): Medical justification supports necessity of the digital component (i.e., rationale why inhaler usage cannot be tracked manually);
5. For requests for Ohtuvayre, both of the following (a and b):
  - a. Member has moderate-to-severe COPD as evidenced by one of the following (i or ii):
    - i. Pre- and post-albuterol forced expiratory volume (FEV<sub>1</sub>)/forced vital capacity (FVC) ratio of  $< 0.70$ ;
    - ii. Post-albuterol FEV<sub>1</sub>  $\geq 30$  % and  $\leq 70$ % of predicted normal;
  - b. Ohtuvayre is not prescribed in combination with Daliresp®;
6. Request meets one of the following (a, b, or c):
  - a. Requested quantity does not exceed the health plan quantity limit;
  - b. Requested dose does not exceed the FDA-approved maximum dose for the relevant indication (see *Section V*);
  - c. Request is for a Georgia member with asthma or other life-threatening bronchial ailments for inhalants prescribed by the treating physician.

**Approval duration: 12 months**

**B. Other diagnoses/indications (must meet 1 or 2):**

1. If this drug has recently (within the last 6 months) undergone a label change (e.g., newly approved indication, age expansion, new dosing regimen) that is not yet reflected in this policy, refer to one of the following policies (a or b):
  - a. For drugs on the formulary (commercial, health insurance marketplace) or PDL (Medicaid), the no coverage criteria policy for the relevant line of business: HIM.PA.33 for health insurance marketplace; or
  - b. For drugs NOT on the formulary (commercial, health insurance marketplace) or PDL (Medicaid), the non-formulary policy for the relevant line of business: HIM.PA.103 for health insurance marketplace; or
2. If the requested use (e.g., diagnosis, age, dosing regimen) is NOT specifically listed under section III (Diagnoses/Indications for which coverage is NOT authorized) AND criterion 1 above does not apply, refer to the off-label use policy for the relevant line of business: HIM.PA.154 for health insurance marketplace.

**II. Continued Therapy**

**A. Inhaled Agents for Asthma or Chronic Obstructive Pulmonary Disease (must meet all):**

1. Member meets one of the following (a or b):
  - a. Currently receiving medication via Centene benefit or member has previously met initial approval criteria;

- b. Member is currently receiving medication and is enrolled in a state and product with continuity of care regulations (*refer to state specific addendums for CC.PHARM.03A and CC.PHARM.03B*);
2. Member is responding positively to therapy;
3. If request is for a dose increase, request meets one of the following (a, b, or c):
  - a. Requested quantity does not exceed the health plan quantity limit;
  - b. Requested dose does not exceed the FDA-approved maximum dose for the relevant indication (see *Section V*);
  - c. Request is for a Georgia member with asthma or other life-threatening bronchial ailments for inhalants prescribed by the treating physician.

**Approval duration: 12 months**

**B. Other diagnoses/indications (must meet 1 or 2):**

1. If this drug has recently (within the last 6 months) undergone a label change (e.g., newly approved indication, age expansion, new dosing regimen) that is not yet reflected in this policy, refer to one of the following policies (a or b):
  - a. For drugs on the formulary (commercial, health insurance marketplace) or PDL (Medicaid), the no coverage criteria policy for the relevant line of business: HIM.PA.33 for health insurance marketplace; or
  - b. For drugs NOT on the formulary (commercial, health insurance marketplace) or PDL (Medicaid), the non-formulary policy for the relevant line of business: HIM.PA.103 for health insurance marketplace; or
2. If the requested use (e.g., diagnosis, age, dosing regimen) is NOT specifically listed under section III (Diagnoses/Indications for which coverage is NOT authorized) AND criterion 1 above does not apply, refer to the off-label use policy for the relevant line of business: HIM.PA.154 for health insurance marketplace.

**III. Diagnoses/Indications for which coverage is NOT authorized:**

- A. Non-FDA approved indications, which are not addressed in this policy, unless there is sufficient documentation of efficacy and safety according to the off label use policies – HIM.PA.154 for health insurance marketplace or evidence of coverage documents.

**IV. Appendices/General Information**

*Appendix A: Abbreviation/Acronym Key*

COPD: chronic obstructive pulmonary disease

EIB: exercise-induced bronchospasm

FDA: Food and Drug Administration

FEV<sub>1</sub>: forced expiratory volume

FVC: forced vital capacity

ICS: inhaled corticosteroid

GINA: Global Initiative for Asthma

GOLD: Global Initiative for Chronic Obstructive Lung Disease

LABA: long acting beta-2 agonist

LAMA: long acting muscarinic antagonist

PDE: phosphodiesterase

SABA: short acting beta-2 agonist

*Appendix B: Therapeutic Alternatives*

*This table provides a listing of preferred alternative therapy recommended in the approval criteria. The drugs listed here may not be a formulary agent for all relevant lines of business and may require prior authorization.*

<b>Drug Name</b>	<b>Dosing Regimen</b>	<b>Dose Limit/ Maximum Dose</b>
fluticasone-salmeterol HFA (Advair HFA authorized generic)	Asthma: 2 inhalations BID (starting dosage is based on asthma severity)	Asthma: 2 inhalations of 230/21 mcg BID
albuterol (Proventil HFA <sup>®</sup> , Ventolin HFA <sup>®</sup> )	<i>Metered-dose inhaler (MDI):</i> 2 puffs every 4 to 6 hours as needed  <i>Nebulization solution:</i> 2.5 mg via oral inhalation every 6 to 8 hours as needed	<i>MDI:</i> 12 puffs/day  <i>Nebulization solution:</i> 4 doses/day or 10 mg/day  Higher maximum dosages for inhalation products have been recommended in National Asthma Education and Prevention Program guidelines for acute exacerbations of asthma.
Anoro Ellipta (umeclidinium/vilanterol)	COPD: 1 inhalation by mouth QD	COPD: 1 inhalation/day
Arcapta Neohaler (indacaterol)	COPD: 75 mcg inhaled orally QD	COPD: 75 mcg/day
Arnuity Ellipta (fluticasone furoate)	Asthma: ≥ 12 years: 100-200 mcg inhaled QD 5-11 years: 50 mcg inhaled QD	Asthma: ≥ 12 years: 200 mcg/day 5-11 years: 50 mcg/day
Breo Ellipta (fluticasone/vilanterol)	Asthma: Age ≥ 18 years: 1 inhalation of 100/25 or 200/25 mcg QD Age 12-17 years: 1 inhalation of 100/25 mcg QD Age 5-11 years: 1 inhalation of 50/25 mcg QD  COPD: 1 inhalation of 100/25 mcg QD	Asthma: 200/25 mcg/day  COPD: 100/25 mcg/day
Breztri Aerosphere	COPD: 2 inhalations by mouth BID	4 inhalations/day
budesonide/formoterol (Symbicort)	Asthma: 2 inhalations BID  COPD: 2 inhalations (160/4.5 mcg) BID	Asthma/COPD: 160/4.5 mcg BID

Drug Name	Dosing Regimen	Dose Limit/ Maximum Dose
Flovent Diskus (fluticasone)	Asthma: 1 inhalation BID (starting dosage is based on asthma severity)	Asthma: 2,000 mcg/day
Flovent HFA (fluticasone)	Asthma: 1 inhalation BID	Asthma: 1,760 mcg/day
fluticasone/salmeterol (Advair Diskus, Wixela Inhub)	Asthma: 1 inhalation BID (starting dosage is based on asthma severity)  COPD: 1 inhalation of 250/50 mcg BID	Asthma: 500/50 mcg BID  COPD: 250/50 mcg BID
Incruse Ellipta (umeclidinium)	COPD: 1 inhalation (62.5 mcg) QD	COPD: 62.5 mcg/day
Pulmicort Flexhaler (budesonide)	Asthma: Starting dose of 180-360 mcg inhaled BID	Asthma: 720 mcg BID
Qvar RediHaler (beclomethasone)	Asthma: ≥ 12 years: 40 mcg, 80 mcg, 160 mcg, or 320 mcg inhaled BID 4-11 years: 40 mcg or 80 mcg inhaled BID	Asthma: ≥ 12 years: 640 mcg/day 4-11 years: 160 mcg/day
Serevent (salmeterol)	Asthma/COPD: 1 inhalation (50 mcg) BID	Asthma/COPD: 100 mcg/day
Spiriva Handihaler (tiotropium bromide monohydrate)	COPD: 2 inhalations (18 mcg) QD	COPD: 18 mcg/day
Spiriva Respimat (tiotropium bromide monohydrate)	Asthma: 2 inhalations (1.25 mcg) QD  COPD: 2 inhalations (2.5 mcg) QD	Asthma: 2.5 mcg/day  COPD: 5 mcg/day
Stiolto Respimat (tiotropium/olodaterol)	Two inhalations by mouth QD at the same time of day	2 inhalations/day
Striverdi Respimat (olodaterol)	COPD: 2 inhalations QD	COPD: 5 mcg/day
Trelegy Ellipta (fluticasone/umeclidinium/vilanterol)	Asthma: 1 inhalation (100/62.5/26 mcg or 200/62.5/26 mcg) by mouth QD  COPD: 1 inhalation (100/62.5/26 mcg) by mouth QD	Asthma: 200/62.5/26 mcg/day  COPD: 100/62.5/26 mcg/day

*Therapeutic alternatives are listed as Brand name<sup>®</sup> (generic) when the drug is available by brand name only and generic (Brand name<sup>®</sup>) when the drug is available by both brand and generic.*

#### Appendix C: Contraindications/Boxed Warnings

- Contraindication(s):
  - All agents: hypersensitivity to any component of the requested agent or the following as additionally specified:
    - Advair Diskus, AirDuo Digihaler/RespiClick, ArmonAir Digihaler, Asmanex Twisthaler, Tudorza Pressair, Trelegy Ellipta, Flovent Diskus: milk proteins

- Brovana: racemic formoterol
- Advair Diskus, AirDuo Digihaler/RespiClick, Alvesco, ArmonAir Digihaler, Asmanex HFA/Twisthaler, Dulera, Pulmicort Respules, Flovent Diskus, Flovent HFA: primary treatment of status asthmaticus or acute episodes of asthma or COPD requiring intensive measures
- Bevespi Aerosphere, Brovana, Duaklir Pressair, Perforomist, Utibron Neohaler: use of a LABA without an ICS in patients with asthma
- Boxed warning(s): none reported

*Appendix D: General Information*

- Although inhaler devices with a digital component may offer increased convenience with tracking of inhaler usage, there is currently no evidence that this leads to improved clinical outcomes, including safety and effectiveness.
- Per the Global Initiative for Chronic Obstructive Lung Disease (GOLD) COPD guidelines, combination therapy (LAMA + LABA or ICS + LAMA + LABA) is recommended for Group B and ED patients (i.e., those who are very symptomatic or are at high risk of exacerbation). Selection of which combination to use depends on the individual patient:
  - For those with more severe symptoms, LAMA + LABA may be used.
  - For those who are inadequately controlled by dual therapy or with blood eosinophil counts at least 300 cells/uL, triple therapy with ICS + LAMA + LABA may be used.
  - As of the 2023 guideline update, use of LABA + ICS in COPD is no longer encouraged. If there is an indication for an ICS, then LABA + LAMA + ICS has been shown to be superior to LABA + ICS and is therefore the preferred choice.
- Historical management of asthma has involved an as-needed short-acting beta agonist for reliever therapy, with stepwise approach to add on controller maintenance therapies such as inhaled corticosteroids and long-acting beta agonists. In 2019, the Global Initiative for Asthma (GINA) guidelines for asthma management and prevention began recommending that inhaled corticosteroids be initiated as soon as possible after diagnosis of asthma, including use as reliever therapy (to be administered as-needed alongside a short-acting beta agonist). The National Asthma Education and Prevention Program from the National Heart, Lung, and Blood Institute followed suit with their recommendations in 2020.
- Alvesco: Use in pediatric patients < 12 years of age: Two identically designed randomized, double-blind, parallel, placebo-controlled clinical trials of 12-weeks treatment duration were conducted in 1,018 patients aged 4 to 11 years with asthma but efficacy was not established. In addition, one randomized, double-blind, parallel, placebo-controlled clinical trial did not establish efficacy in 992 patients aged 2 to 6 years with asthma.

**V. Dosage and Administration**

<b>Drug Name</b>	<b>Indication</b>	<b>Dosing Regimen</b>	<b>Maximum Dose</b>
Advair Diskus	Asthma	1 inhalation BID (starting dosage is based on asthma severity)	500/50 mcg BID
	COPD	1 inhalation of 250/50 mcg BID	250/50 mcg BID
Advair HFA	Asthma	2 inhalations BID (starting dosage is based on asthma severity)	2 inhalations of 230/21 mcg BID



<b>Drug Name</b>	<b>Indication</b>	<b>Dosing Regimen</b>	<b>Maximum Dose</b>
AirDuo Digihaler	Asthma	1 inhalation BID (starting dosage is based on asthma severity)	232/14 mcg BID
AirDuo RespiClick	Asthma	1 inhalation BID (starting dosage is based on asthma severity)	232/14 mcg BID
Alvesco	Asthma	Starting dose for patients who received bronchodilators alone: 80 mcg inhaled BID  Starting dose for patients who received inhaled corticosteroids: 80 mcg inhaled BID  Starting dose for patients who received oral corticosteroids: 320 mcg inhaled BID	320 mcg/day  640 mcg/day  640 mcg/day
ArmonAir Digihaler	Asthma	1 inhalation BID (starting dosage is based on asthma severity and age)	232 mcg BID
Asmanex HFA	Asthma	2 inhalations BID (starting dosage is based on age and asthma severity)	800 mcg/day
Asmanex Twisthaler	Asthma	Dose varies based on previous therapy and age: 1 inhalation QD-BID	880 mcg/day
Bevespi Aerosphere	COPD	2 inhalations BID	4 inhalations/day
Brovana	COPD	One 15 mcg/2 mL vial inhaled via nebulizer every 12 hours	30 mcg/day
Duaklir Pressair	COPD	One inhalation by mouth BID	2 inhalations/day
Dulera	Asthma	Age 5 to 11 years: 2 inhalations of 50/5 mcg BID Age ≥ 12 years: 2 inhalations of 100/5 mcg or 200/5 mcg BID (starting dosage is based on asthma severity)	200/5 mcg/day 800/20 mcg/day
Flovent Diskus	Asthma	1 inhalation BID (starting dosage is based on asthma severity)	1,000 mcg BID
Flovent HFA	Asthma	Patients aged 12 years and older: 88 mcg twice daily up to a maximum dosage of 880 mcg twice daily. Pediatric patients aged 4 to 11 years: 88 mcg twice daily	880 mcg BID
Lonhala Magnair	COPD	One 25 mcg vial inhaled via nebulizer BID	50 mcg/day
Ohtuvayre	COPD	3 mg (one ampule) inhaled via nebulizer BID	6 mg/day

Drug Name	Indication	Dosing Regimen	Maximum Dose
Perforomist	COPD	One 20 mcg/2 mL vial inhaled via nebulizer every 12 hours	40 mcg/day
ProAir Digihaler	Treatment or prevention of bronchospasm	2 inhalations every 4 to 6 hours	12 inhalations/day
	Prevention of EIB	2 inhalations 15 to 30 minutes before exercise	2 inhalations before exercise
Pulmicort Respules	Asthma	Starting dose for patients who received bronchodilators alone or inhaled corticosteroids: 0.5 mg inhaled per day (0.5 mg QD or 0.25 mg BID; for inhaled corticosteroids, may go up to 0.5 mg BID)  Starting dose for patients who received oral corticosteroids: 1 mg inhaled per day (1 mg QD or 0.5 mg BID)	Bronchodilator alone: 0.5 mg/day  Inhaled or oral corticosteroid: 1 mg/day
Seebri Neohaler	COPD	One inhalation (15.6 mcg) BID	2 inhalations/day
Symbicort	Asthma	2 inhalations BID (starting dosage is based on asthma severity)	320/9 mcg BID
	COPD	2 inhalations (160/4.5 mcg) BID	320/9 mcg BID
Symbicort Aerosphere	COPD	2 inhalations (160/4.8 mcg) BID	320/9.6 mcg BID
Tudorza Pressair	COPD	1 inhalation (400 mcg) BID	800 mcg/day
Utibron Neohaler	COPD	Inhalation of the contents of one capsule BID	2 capsules/day
Yupelri	COPD	One 175 mcg mcg vial inhaled via nebulizer QD	175 mcg/day

## VI. Product Availability

Drug Name	Availability
Advair Diskus	Inhalation powder containing fluticasone/salmeterol: 100/50 mcg, 250/50 mcg, 500/50 mcg
Advair HFA	Inhalation aerosol containing fluticasone/salmeterol: 45/21 mcg, 115/21 mcg, 230/21 mcg
AirDuo Digihaler	Inhalation powder: In each actuation: 55/14 mcg contains 55 mcg of fluticasone propionate and 14 mcg of salmeterol; 113/14 mcg contains 113 mcg of fluticasone propionate and 14 mcg of salmeterol; 232/14 mcg contains 232 mcg of fluticasone propionate and 14 mcg of salmeterol. AirDuo Digihaler contains a built-in electronic module
AirDuo RespiClick	Inhalation powder: In each actuation: 55 mcg/14 mcg contains 55 mcg of fluticasone propionate and 14 mcg of salmeterol; 113 mcg/14 mcg

Drug Name	Availability
	contains 113 mcg of fluticasone propionate and 14 mcg of salmeterol; 232 mcg/14 mcg contains 232 mcg of fluticasone propionate and 14 mcg of salmeterol
Alvesco	Inhalation aerosol: 80 mcg/actuation, 160 mcg/actuation
ArmonAir Digihaler	Inhalation powder containing 30 mcg, 55 mcg, 113 mcg, or 232 mcg of fluticasone propionate per actuation. ArmonAir Digihaler contains a built-in electronic module
Asmanex HFA	Inhalation aerosol containing 50 mcg, 100 mcg, or 200 mcg of mometasone furoate per actuation
Asmanex Twisthaler	Inhalation device: 110 mcg (delivers 100 mcg/actuation), 220 mcg (delivers 200 mcg/actuation)
Bevespi Aerosphere	Inhalation aerosol: pressurized metered dose inhaler containing a combination of glycopyrrolate (9 mcg) and formoterol fumarate (4.8 mcg) per inhalation; two inhalations equal one dose
Brovana	Inhalation solution (unit-dose vial for nebulization): 15 mcg/2 mL
Duaklir Pressair	Inhalation powder: 30 and 60 metered dose dry powder inhaler metering 400 mcg aclidinium bromide and 12 mcg formoterol fumarate per actuation
Dulera	Inhalation aerosol containing mometasone/formoterol: 50/5 mcg, 100/5 mcg, 200/5 mcg per actuation
Flovent Diskus	Inhalation powder: Inhaler containing fluticasone propionate (50, 100, or 250 mcg) as a powder formulation for oral inhalation
Flovent HFA	Inhalation aerosol: 44 mcg, 110 mcg, 220 mcg per actuation
Lonhala Magnair	Sterile solution for inhalation in a unit-dose vial: 25 mcg/mL
Ohtuvayre	Inhalation suspension in unit-dose ampule: 3 mg/2.5 mL
Perforomist	Inhalation solution (unit dose vial for nebulization): 20 mcg/2 mL solution
ProAir Digihaler	Inhalation powder: dry powder inhaler 108 mcg of albuterol sulfate (equivalent to 90 mcg of albuterol base) from the mouthpiece per actuation. The inhaler is supplied for 200 inhalation doses. ProAir Digihaler includes a built-in electronic module
Pulmicort Respules	Inhalation suspension: 0.25 mg/2 mL, 0.5 mg/2 mL, 1 mg/2 mL
Seebri Neohaler	Inhalation powder in capsules: 15.6 mcg of glycopyrrolate inhalation powder for use with the Neohaler device
Symbicort	Metered-dose inhaler: budesonide (80 or 160 mcg) and formoterol (4.5 mcg) as an inhalation aerosol
Symbicort Aerosphere	Metered-dose inhaler: budesonide (160 mcg) and formoterol (4.8 mcg) as an inhalation aerosol
Tudorza Pressair	Inhalation powder in a multi-dose dry powder inhaler: 400 mcg/actuation
Utibron Neohaler	Inhalation powder in capsule, for use with the Neohaler device: 27.5 mcg of indacaterol and 15.6 mcg glycopyrrolate
Yupelri	Inhalation solution (unit-dose vial for nebulization): 175 mcg/3 mL

## VII. References

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Reviews, Revisions, and Approvals	Date	P&T Approval Date
Policy created: adapted from previously approved individual drug policies- CP.PCH.35 Alvesco, CP.PCH.36 Asmanex, HIM.PA.48 Pulmicort Respules, HIM.PA.102 Utibron Neohaler, HIM.PA.150 Breztri Aerosphere, and HIM.PA.151 Duaklir Pressair (all to be retired); added additional agents and revised criteria to reflect SDC CY2021 strategy/prior clinical guidance; added requirement for medical justification for requests for agents with digital component.	10.29.20	02.21
Added option for request to not exceed the health plan quantity limit.	04.23.21	
Per October SDC, removed Breztri Aerosphere from criteria.	10.27.21	
1Q 2022 annual review: no significant changes; references reviewed and updated.	09.21.21	02.22
Per March SDC for brand Perforomist and Brovana added redirection to generic formoterol/arformoterol.	03.22.22	05.22
RT4: updated ArmonAir Digihaler per prescribing information for pediatric extension down to 4 years of age and older, added new 30 mcg strength; references reviewed and updated.	05.16.22	
Per August SDC, revised AirDuo Digihaler, AirDuo RespiClick, Dulera redirection to include only Symbicort authorized generic rather than both brand and generic. Template changes applied to other diagnoses/indications and continued therapy section.	08.23.22	11.22
1Q 2023 annual review: no significant changes; updated Appendix D with updated 2023 GOLD guideline recommendations; references reviewed and updated.	01.11.23	02.23
Per April SDC, removed Xopenex from policy.	04.20.23	
RT4: added newly approved dosage form Symbicort Aerosphere to policy with redirection to generic Symbicort per SDC and prior clinical guidance; updated dosing for Breo Ellipta in Appendix B per prescribing information for pediatric extension down to 5 years of age and older. Corrected maximum dose for Bevespi Aerosphere from 2 inhalations/day to 4 inhalations/day per dosing regimen (2 inhalations BID); added redirection to generic Symbicort for brand Symbicort per SDC and prior clinical guidance.	05.26.23	
Per April SDC and prior clinical guidance, revised redirection from brand Flovent HFA/Flovent Diskus to instead redirect to fluticasone propionate HFA (Flovent HFA authorized generic); added Flovent HFA and Advair HFA to policy requiring redirection to authorized generic; revised redirection to brand Advair HFA to instead redirect to authorized generic; for LABA/LAMA revised redirection to Bevespi Aerosphere to instead redirect to Stiolto Respimat; added Bevespi Aerosphere to policy with redirection to Anoro Ellipta and Stiolto Respimat; for AirDuo Digihaler, AirDuo RespiClick, Dulera, updated redirection to include both Breo Ellipta authorized generic	09.21.23	12.23

Reviews, Revisions, and Approvals	Date	P&T Approval Date
and brand Breo Ellipta. Added dose/quantity limit bypass for Georgia members with asthma or other life-threatening bronchial ailments for inhalants prescribed by the treating physician.		
1Q 2024 annual review: no significant changes; references reviewed and updated.	10.09.23	02.24
Per March SDC, for “All other ICS” requests added additional redirection to fluticasone propionate diskus (Flovent Diskus authorized generic); revised Flovent Diskus redirection requirements to fluticasone propionate diskus (Flovent Diskus authorized generic) in a new row; for “All other ICS/LABA” requests added additional redirection to Dulera.	03.12.24	05.24
RT4: added newly approved agent Ohtuvayre with redirections per SDC.	07.03.24	11.24

**Important Reminder**

This clinical policy has been developed by appropriately experienced and licensed health care professionals based on a review and consideration of currently available generally accepted standards of medical practice; peer-reviewed medical literature; government agency/program approval status; evidence-based guidelines and positions of leading national health professional organizations; views of physicians practicing in relevant clinical areas affected by this clinical policy; and other available clinical information. The Health Plan makes no representations and accepts no liability with respect to the content of any external information used or relied upon in developing this clinical policy. This clinical policy is consistent with standards of medical practice current at the time that this clinical policy was approved. “Health Plan” means a health plan that has adopted this clinical policy and that is operated or administered, in whole or in part, by Centene Management Company, LLC, or any of such health plan’s affiliates, as applicable.

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This clinical policy is effective as of the date determined by the Health Plan. The date of posting may not be the effective date of this clinical policy. This clinical policy may be subject to applicable legal and regulatory requirements relating to provider notification. If there is a discrepancy between the effective date of this clinical policy and any applicable legal or regulatory requirement, the requirements of law and regulation shall govern. The Health Plan retains the right to change, amend or withdraw this clinical policy, and additional clinical policies may be developed and adopted as needed, at any time.



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