



Prior Authorization Request Administrative Information

Member information

Last name First name MI

Member ID Date of birth

Sex assigned at birth Female Male "X" or Intersex

Current gender Female Male Transgender male Transgender female Other

Place of residence Home Nursing facility Other

Race Ethnicity

Preferred spoken language Preferred written language

MassHealth does not exclude people or treat them differently because of race, color, national origin, age, disability, religion, creed, sexual orientation, or sex (including gender identity and gender stereotyping).

Plan contact information

Please indicate the member's MassHealth Plan and fax or submit this completed and signed form according to the Plan's contact information below.

MassHealth Fee-For-Service (FFS) Plan, Primary Care Clinician (PCC) Plan, Primary Care Accountable Care Organization (PCACO) Plan, and Children's Medical Security Plan
<input type="checkbox"/> MassHealth Drug Utilization Review Program Pharmacy: Fax: (877) 208-7428 - Tel: (800) 745-7318
MassHealth Managed Care Organization (MCO) and Accountable Care Partnership Plans (ACPP)
<input type="checkbox"/> Fallon Health Online Prior Authorization: go.covermymeds.com/OptumRx Online Prior Authorization: providerportal.surescripts.net/ProviderPortal/optum Pharmacy: Fax: (844) 403-1029 - Tel: (844) 720-0033
<input type="checkbox"/> Health New England Online Prior Authorization: go.covermymeds.com/OptumRx Pharmacy: Fax: (800) 550-9246 - Tel: (800) 918-7545
<input type="checkbox"/> Mass General Brigham Health Plan Online Prior Authorization (Pharmacy Benefit Reviews): go.covermymeds.com/OptumRx Pharmacy: Fax: (844) 403-1029 - Tel: (800) 711-4555 Online Prior Authorization (Medical Specialty Reviews): provider.massgeneralbrighamhealthplan.org Medical Specialty Reviews: Fax: (888) 656-6671 - Tel: (833) 895-2611
<input type="checkbox"/> Tufts Health Plan Online Prior Authorization: point32health.promptpa.com Pharmacy: Fax: (617) 673-0939 - Tel: (888) 257-1985
<input type="checkbox"/> WellSense Health Plan Online Prior Authorization: wellsense.org/providers/ma/pharmacy/prior-authorizations Pharmacy: Fax: (833) 951-1680 - Tel: (877) 417-1822

Inhaled Respiratory Agents Prior Authorization Request

MassHealth reviews requests for prior authorization (PA) on the basis of medical necessity only. If MassHealth approves the request, payment is still subject to all general conditions of MassHealth, including current member eligibility, other insurance, and program restrictions. MassHealth will notify the requesting provider and member of its decision. Keep a copy of this form for your records. If faxing this form, please use black ink.

Additional information about these agents, including PA requirements and preferred products, can be found within the MassHealth Drug List at www.mass.gov/druglist.

Medication information

Medication requested (Check one or all that apply. Where applicable, the brand name is provided in brackets for reference.)

Anticholinergics

- Yupelri (revefenacin)
- Tudorza (aclidinium)

Combination Products

- Airsupra (albuterol/budesonide)
- Bevespi (glycopyrrolate/formoterol)
- Breztri (budesonide/glycopyrrolate/formoterol)
- Duaklir (aclidinium/formoterol)
- fluticasone/salmeterol [Airduo Respiclick]
- Stiolto (tiotropium/olodaterol)
- Trelegy (fluticasone furoate/umeclidinium/vilanterol)

Corticosteroids

- Alvesco (ciclesonide inhaler)
- budesonide inhalation suspension \geq 13 years
- fluticasone propionate inhalation aerosol \geq 12 years
- fluticasone propionate inhalation powder
- Qvar Redihaler (beclomethasone inhaler)

Long-acting Beta Agonists

- arformoterol

- formoterol
- Striverdi (olodaterol)

Short-acting Beta Agonists

- albuterol inhaler ‡
- levalbuterol inhalation solution

‡Brand name Ventolin is available without prior authorization.

Phosphodiesterase 3/phosphodiesterase 4 inhibitor

- Ohtuvayre (ensifentrine)

Other Medication

- Other*

*If request is for a non-preferred brand name or generic product, please attach supporting documentation (e.g., copies of medical records and/or office notes regarding adverse reaction or inadequate response to the preferred product).

Dose and frequency of medication requested

Number of inhalers/month

Indication (Check all that apply or include ICD-10 code, if applicable.)

- Asthma (Specify severity below.)
 - Intermittent
 - Mild Persistent
 - Moderate Persistent
 - Severe Persistent
- Chronic Obstructive Pulmonary Disease (COPD) (Specify severity and subtype below.)
 - Severity Mild Moderate Severe Very severe
 - Subtype Chronic bronchitis Emphysema

Exercise-induced bronchospasm

Reactive airway disease

Other

Please list all other medications currently prescribed for the member for this indication.

Is this member a referral candidate for care coordination? Yes No

If yes, MassHealth will offer care coordination services to this member. Please describe which additional behavioral health services would be beneficial. *Please inform the member, parent, or legal guardian to expect outreach from a MassHealth representative of care coordination services.*

Section I. Please complete for albuterol inhaler requests.

1. For requests for albuterol inhaler, has the member had a trial with an albuterol product available without prior authorization? *

Yes. Please list the dates/duration of trials, and outcomes in Section XI.

No. Please describe the clinical rationale why albuterol product available without prior authorization is not appropriate for this member.

* Brand name Ventolin does not require prior authorization.

Section II. Please complete for all arformoterol, budesonide inhalation suspension, formoterol, levalbuterol inhalation solution, and Yupelri requests.

1. Please describe the medical necessity for a nebulized formulation.

2. For levalbuterol inhalation solution, has the member had a trial with albuterol solution?

Yes. Please list the dates/duration of trials, and outcomes in Section XI.

No. Please describe the clinical rationale why albuterol solution is not appropriate for this member.

3. For Yupelri, has the member had a trial with ipratropium inhalation nebulizer solution?

Yes. Please list the dates/duration of trials, and outcomes in Section XI.

No. Please describe the clinical rationale why ipratropium inhalation nebulizer solution is not appropriate

for this member.

Section III. Please complete for fluticasone/salmeterol (generic Airduo Resplick) requests.

1. Has the member had a trial with fluticasone/salmeterol inhalation aerosol, powder (generic Advair)?

Yes. Please list the dates/duration of trials and the outcomes in Section XI.

No. Please describe the clinical rationale for use of the requested agent in this member.

Section IV. Please complete for Alvesco, fluticasone propionate inhalation aerosol for members ≥ 12 years of age, fluticasone propionate inhalation powder, and Qvar Redihaler requests.

Has the member had a trial with two inhaled corticosteroids?

Yes. Please list the dates/duration of trials, and outcomes in Section XI.

No. Please document if there is a contraindication to all other inhaled corticosteroids.

Section V. Please complete for Bevespi and Duaklir requests.

Has the member had a trial with Stiolto or umeclidinium/vilanterol?

Yes. Please list the dates/duration of trials, and outcomes in Section XI.

No. Please describe the clinical rationale for use of the requested agent in this member.

Section VI. Please complete for Trelegy requests.

Has the member had a trial with fluticasone/vilanterol and umeclidinium or fluticasone furoate inhalation powder and umeclidinium/vilanterol?

Yes. Please list the dates/duration of trials, and outcomes in Section XI.

No. Please describe the clinical rationale for use of the requested agent in this member.

Section VII. Please complete for Breztri requests.

Has the member had a trial with the following combination of the separate agents: Bevespi and Pulmicort inhalation powder?

Yes. Please list the dates/duration of trials, and outcomes in Section XI.

No. Please describe the clinical rationale for use of the requested agent in this member.

Section VIII. Please complete for Airsupra requests.

Has the member had a trial with budesonide/formoterol or albuterol and Pulmicort inhalation powder?

Yes. Please list the dates/duration of trials, and outcomes in Section XI.

No. Please describe the clinical rationale for use of the requested agent in this member.

Section IX. Please complete for Ohtuvayre requests.

1. Has the member had a trial with Bevespi, Duaklir, Stiolto, or umeclidinium/vilanterol?

Yes. Please list the dates/duration of trials, and outcomes in Section XI.

No. Please describe the clinical rationale why Bevespi, Duaklir, Stiolto, and umeclidinium/vilanterol is not appropriate for this member.

2. Has the member had a trial with Breztri or Trelegy?

Yes. Please list the dates/duration of trials, and outcomes in Section XI.

No. Please describe the clinical rationale why Breztri and Trelegy is not appropriate for this member.

Section X. Please complete for Tudorza requests.

1. Has the member had a trial with Spiriva Respimat, tiotropium inhalation powder, or umeclidinium?

Yes. Please list the dates/duration of trials, and outcomes in Section X.

No. Please document if there is a contraindication to Spiriva Respimat, tiotropium inhalation powder, and umeclidinium.

Section XI. Please complete as instructed in sections above.*

Drug name Dates/duration of use

Did the member experience any of the following? Adverse reaction Inadequate response Other

Briefly describe details of adverse reaction, inadequate response, or other.

Drug name Dates/duration of use

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Briefly describe details of adverse reaction, inadequate response, or other.

Drug name Dates/duration of use

Did the member experience any of the following? Adverse reaction Inadequate response Other

Briefly describe details of adverse reaction, inadequate response, or other.

** Please attach a letter documenting additional trials as necessary.*

Section XII. Please complete and provide documentation for exceptions to step therapy.

1. Is the alternative drug required under the step therapy protocol contraindicated, or will likely cause an adverse reaction in, or physical or mental harm to, the member? Yes No

If yes, briefly describe details of contraindication, adverse reaction, or harm.

2. Is the alternative drug required under the step therapy protocol expected to be ineffective based on the known clinical characteristics of the member and the known characteristics of the alternative drug regimen? Yes No

If yes, briefly describe details of known clinical characteristics of member and alternative drug regimen.

3. Has the member previously tried the alternative drug required under the step therapy protocol, or another alternative drug in the same pharmacologic class or with the same mechanism of action, and such alternative drug was discontinued due to lack of efficacy or effectiveness, diminished effect, or an adverse event? Yes No

If yes, please provide details for the previous trial.

Drug name Dates/duration of use

Did the member experience any of the following? Adverse reaction Inadequate response

Briefly describe details of adverse reaction or inadequate response.

4. Is the member stable on the requested prescription drug prescribed by the healthcare provider, and switching drugs will likely cause an adverse reaction in, or physical or mental harm to, the member?

Yes. Please provide details.

No

Please continue to next page and complete Prescriber and Provider Information section.

Prior Authorization Request Prescriber and Provider Information

Prescriber information

Last name*	<input type="text"/>	First name*	<input type="text"/>	MI	<input type="text"/>
NPI*	<input type="text"/>	Individual MH Provider ID	<input type="text"/>		
DEA No.	<input type="text"/>	Office Contact Name	<input type="text"/>		
Address	<input type="text"/>	City	<input type="text"/>	State	<input type="text"/>
		Zip	<input type="text"/>		
E-mail address	<input type="text"/>				
Telephone No.*	<input type="text"/>				
Fax No.* (Please provide fax number for PA response notification.)	<input type="text"/>				

* Required

Please also complete for professionally administered medications, if applicable.

Start date	<input type="text"/>	End date	<input type="text"/>		
Servicing prescriber/facility name	<input type="text"/>	<input type="checkbox"/>	Same as prescribing provider		
Servicing provider/facility address	<input type="text"/>				
Servicing provider NPI/tax ID No.	<input type="text"/>				
Name of billing provider	<input type="text"/>				
Billing provider NPI No.	<input type="text"/>				
Is this a request for recertification?	<input type="checkbox"/>	Yes	<input type="checkbox"/>	No	
CPT code	<input type="text"/>	No. of visits	<input type="text"/>	J code	<input type="text"/>
				No. of units	<input type="text"/>

Provider's attestation, signature, and date

I certify under the pains and penalties of perjury that I am either the prescribing provider or duly authorized to act on behalf of the provider identified in the Prescriber information section of this form. Any attached statement on letterhead has been reviewed and signed by me. I certify that the medical necessity information (per 130 CMR 450.204) on this form is true, accurate, and complete, to the best of my knowledge. I understand that I may be subject to civil penalties or criminal prosecution for any falsification, omission, or concealment of any material fact contained herein.

Signature of provider or individual duly authorized to act on behalf of the provider:

Printed legal name and title of signatory above

<input type="text"/>	Date	<input type="text"/>
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(The form can either be signed by hand and then scanned, or it can be signed electronically using DocuSign or Adobe Sign. For electronic signatures, the signer can upload a picture of their wet signature. The typed text of a signature is not an acceptable form of an electronic signature.)