



Commonwealth of Massachusetts
MassHealth Drug Utilization Review Program
 P.O. Box 2586, Worcester, MA 01613-2586
Fax: (877) 208-7428 **Phone:** (800) 745-7318

Antihistamine 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.

Member information

Last name _____ First name _____ MI _____
 MassHealth member ID # _____ Date of birth _____
 Gender (Check one.) F M Member's place of residence home nursing facility

Medication information

Medication requested (Check one or all that apply.)

Intranasal Antihistamines

azelastine 0.15% nasal spray olopatadine nasal spray

Single Oral Antihistamines

carbinoxamine 6 mg tablet Karbinal ER (carbinoxamine extended-release)
 desloratadine orally disintegrating tablet (ODT) levocetirizine solution
 dexchlorpheniramine solution

Combination Oral Antihistamines

Clarinex-D (desloratadine/pseudoephedrine) Semprex-D (acrivastine/pseudoephedrine)

Injectable Antihistamines

Quzyttir (cetirizine injection)^

Dose and frequency of medication requested _____

^This drug is available through the health care professional who administers the drug. MassHealth does not pay for this drug to be dispensed through a retail pharmacy.

Indication (Check all that apply.)

Allergic Rhinitis Urticaria
 Non-allergic Rhinitis Other _____

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

Section I. Please complete for desloratadine ODT and levocetirizine solution requests. Please also complete Section II for these medications.

Does the member have a swallowing disorder or condition affecting swallowing ability?

- Yes. Please describe. _____
- No. Please describe clinical rationale for not using oral tablet formulation. _____
-

Section II. Please complete for Clarinex-D, desloratadine ODT, levocetirizine solution, and Semprex-D requests.

Has the member had a trial with an intranasal corticosteroid and two second-generation antihistamines (e.g., cetirizine, desloratadine, levocetirizine, loratadine)?

For requests for combination antihistamines, please include information about trials with second-generation antihistamines in combination with pseudoephedrine (e.g., cetirizine/pseudoephedrine, loratadine/pseudoephedrine).

- Yes. Please list the drug names, dates/duration of trials, and outcomes below.*

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 _____

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 _____

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

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

- No. Please describe why intranasal corticosteroids and second-generation antihistamines are not appropriate for this member. _____
-

Section III. Please complete for azelastine 0.15% nasal spray and olopatadine nasal spray requests.

1. Has the member had a trial with an intranasal corticosteroid?

- Yes. Please list the drug name, dates/duration of trials, and outcomes below.*

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.

- No. Please describe why intranasal corticosteroids are not appropriate for this member.
-

2. Has the member had a trial with azelastine 137 mcg nasal spray?

- Yes. Please list the dates/duration of trials and outcomes below.*

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.

- No. Please describe why azelastine 137 mcg nasal spray is not appropriate for this member.
-

3. For the diagnosis of allergic rhinitis, has the member had a trial with two second-generation antihistamines (e.g., cetirizine, desloratadine, levocetirizine, loratadine)?

Yes. Please list the drug names, dates/duration of trials, and outcomes below.*

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 _____

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

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

No. Please describe why second-generation antihistamines are not appropriate for this member.

4. For requests for any agent at a quantity > 1 inhaler per month, please document an inadequate response to the manufacturer's recommended dosing.

Section IV. Please complete for carbinoxamine 6 mg tablet and Karbinal ER requests.

1. Has the member had a trial with an intranasal corticosteroid and two non-selective antihistamines?

Yes. Please list the drug names, dates/duration of trials, and outcomes below.*

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 _____

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 _____

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

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

No. Please describe why intranasal corticosteroids and non-selective antihistamines are not appropriate for this member.

2. For carbinoxamine 6 mg tablet requests, please provide clinical rationale for use instead of carbinoxamine 4 mg tablet and immediate-release carbinoxamine solution.

3. For Karbinal ER requests, has the member had a trial with carbinoxamine immediate-release solution?

Yes. Please list the dates/duration of trials and outcomes below.*

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.

No. Please describe why carbinoxamine immediate-release solution is not appropriate for this member.

*Please attach a letter documenting additional trials as necessary.

Section V. Please complete for dexchlorpheniramine solution requests.

Please provide medical necessity for use of the requested agent instead of all other first- and second-generation antihistamines available in liquid formulation.

Section VI. Please complete for Quzyttir requests.

Has the member had a trial with injectable diphenhydramine?

Yes. Please list the drug names, dates/duration of trials, and outcomes below.

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.

No. Please describe why injectable diphenhydramine is not appropriate for this member.

Section VII. Please complete for all requests for non-preferred drug products if one or more preferred drug products have been designated for this class of drugs.

If one or more preferred drug products have been designated for this class of drugs, and if you are requesting PA for a non-preferred drug product, please provide medical necessity for prescribing the non-preferred drug product rather than the preferred drug product.

Prescriber information

Last name* _____ First name* _____ MI _____

NPI* _____ Individual MH Provider ID _____

DEA No. _____ Office Contact Name _____

Address _____ City _____ State _____ Zip _____

E-mail address _____

Telephone No.* _____ Fax No.* _____

* *Required*

Prescribing provider's attestation, signature, and date

I certify under the pains and penalties of perjury that I am the prescribing provider identified in the Prescriber information section of this form. Any attached statement on my 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.

Prescribing provider's signature (Signature and date stamps, or the signature of anyone other than the provider, are not acceptable.)

Signature required _____

Printed name of prescribing provider _____ Date _____