



Enjaymo

Prior Authorization Request

CVS Caremark administers the prescription benefit plan for the patient identified. This patient's benefit plan requires prior authorization for certain medications in order for the drug to be covered. To make an appropriate determination, providing the most accurate diagnosis for the use of the prescribed medication is necessary. **Please respond below and fax this form to CVS Caremark toll-free at 1-855-330-1720.** If you have questions regarding the prior authorization, please contact CVS Caremark at **1-888-877-0518**. For inquiries or questions related to the patient's eligibility, drug copay or medication delivery; please contact the Specialty Customer Care Team: CaremarkConnect® 1-800-237-2767.

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Patient's Name: _____ **Date:** _____
Patient's ID: _____ **Patient's Date of Birth:** _____
Physician's Name: _____
Specialty: _____ **NPI#:** _____
Physician Office Telephone: _____ **Physician Office Fax:** _____

Referring Provider Info: Same as Requesting Provider

Name: _____ **NPI#:** _____
Fax: _____ **Phone:** _____

Rendering Provider Info: Same as Referring Provider Same as Requesting Provider

Name: _____ **NPI#:** _____
Fax: _____ **Phone:** _____

Approvals may be subject to dosing limits in accordance with FDA-approved labeling, accepted compendia, and/or evidence-based practice guidelines.

Required Demographic Information:

Patient Weight: _____ *kg*

Patient Height: _____ *cm*

Send completed form to: Case Review Unit CVS Caremark Specialty Programs Fax: 1-855-330-1720

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Site of Service Questions:

- A. Where will this drug be administered?
 Ambulatory surgical, *skip to Clinical Questions* Home infusion, *skip to Clinical Questions*
 Off-campus Outpatient Hospital On-campus Outpatient Hospital
 Physician office, *skip to Clinical Questions* Pharmacy, *skip to Clinical Questions*
- B. Is this request to continue previously established treatment with the requested medication?
 Yes - This is a continuation of an existing treatment.
 No - This is a new therapy request (patient has not received requested medication in the last 6 months). *skip to Clinical Criteria Questions*
- C. Has the patient experienced an adverse event with the requested product that has not responded to conventional interventions (eg acetaminophen, steroids, diphenhydramine, fluids, other pre-medications or slowing of the infusion rate) or a severe adverse event (anaphylaxis, anaphylactoid reactions, myocardial infarction, thromboembolism, or seizures) during or immediately after an infusion? ***ACTION REQUIRED: If Yes, Attach supporting clinical documentation.*** Yes, *skip to Clinical Criteria Questions* No
- D. Is the patient medically unstable which may include respiratory, cardiovascular, or renal conditions that may limit the member's ability to tolerate a large volume or load or predispose the member to a severe adverse event that cannot be managed in an alternate setting without appropriate medical personnel and equipment? ***ACTION REQUIRED: If Yes, Attach supporting clinical documentation.*** Yes, *skip to Clinical Criteria Questions* No
- E. Does the patient have significant behavioral issues and/or physical or cognitive impairment that would impact the safety of the infusion therapy AND the patient does not have access to a caregiver? ***ACTION REQUIRED: If Yes, Attach supporting clinical documentation.*** Yes No

Clinical Criteria Questions:

1. What is the diagnosis?
 Cold Agglutinin disease (CAD) Other, please specify _____
2. What is the ICD-10 code? _____
3. Is the request for continuation of therapy with the requested medication? Yes No *If No, skip to #6*
4. Has the patient experienced disease progression or unacceptable toxicity while on the current regimen?
 Yes No
5. Has the patient demonstrated a positive response to therapy (e.g., improvement in hemoglobin levels, improvement in markers of hemolysis [e.g., bilirubin, haptoglobin, lactate dehydrogenase [LDH], reticulocyte count], a reduction in blood transfusions)? ***ACTION REQUIRED: If 'Yes', supporting chart notes documenting a positive response to therapy (e.g., improvement in hemoglobin levels, improvement in markers of hemolysis [e.g., bilirubin, haptoglobin, lactate dehydrogenase [LDH], reticulocyte count], a reduction in blood transfusions) are required***
 Yes No *No further questions.*
6. Was the diagnosis of primary cold agglutinin disease (CAD) confirmed by evidence of hemolysis? Yes No
7. Does the patient have a lactate dehydrogenase (LDH) level above the upper limit of normal? ***ACTION REQUIRED: If Yes, attach chart notes, medical records or test results supporting hemolysis result.***
 Yes No
8. Does the patient have a haptoglobin level below the lower limit of normal? ***ACTION REQUIRED: If Yes, attach chart notes, medical records or test results supporting hemolysis result.*** Yes No
9. Does the patient have a positive polyspecific direct antiglobulin test (DAT) result? ***ACTION REQUIRED:***

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If Yes, attach chart notes, medical records or test results supporting polyspecific direct antiglobulin test (DAT) result. Yes No

10. Does the patient have a monospecific direct antiglobulin test (DAT) result strongly positive for C3d? ***ACTION REQUIRED: If Yes, attach chart notes, medical records or test results supporting monospecific direct antiglobulin test (DAT) result.*** Yes No
11. Does the patient have a cold agglutinin titer of 1:64 or higher measured at 4°C ? ***ACTION REQUIRED: If Yes, attach chart notes, medical records or test results supporting cold agglutinin titer measured at 4°C.*** Yes No
12. Does the patient have a DAT IgG level of 1+ or less? ***ACTION REQUIRED: If Yes, attach chart notes, medical records or test results supporting IgG level.*** Yes No
14. Has secondary cold agglutinin disease (CAD) been ruled out for the patient (e.g., cold agglutinin syndrome secondary to infection, rheumatologic disease, or active hematologic malignancy)? ***ACTION REQUIRED: Please attach chart notes, medical records or test results (e.g., bone marrow biopsy, imaging) ruling out secondary cold agglutinin (CAD).*** Yes No

I attest that this information is accurate and true, and that documentation supporting this information is available for review if requested by CVS Caremark or the benefit plan sponsor.

X _____

Prescriber or Authorized Signature

Date (mm/dd/yy)

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