



Enjaymo[®] (sutimlimab-jome) Medication Precertification Request

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(All fields must be completed and legible for precertification review.)

Aetna Precertification Notification

Phone: 1-866-752-7021

FAX: 1-888-267-3277

For Medicare Advantage Part B:

Please Use Medicare Request Form

Please indicate: Start of treatment: Start date ____ / ____ / ____
 Continuation of therapy, Date of last treatment ____ / ____ / ____

Precertification Requested By: _____ Phone: _____ Fax: _____

A. PATIENT INFORMATION

First Name:		Last Name:		DOB:	
Address:		City:		State:	ZIP:
Home Phone:	Work Phone:	Cell Phone:		Email:	
Patient Current Weight: ____ lbs or ____ kgs				Patient Height: ____ inches or ____ cms	
Allergies:					

B. INSURANCE INFORMATION

Aetna Member ID #: _____	Does patient have other coverage? <input type="checkbox"/> Yes <input type="checkbox"/> No
Group #: _____	If yes, provide ID#: _____ Carrier Name: _____
Insured: _____	Insured: _____
Medicare: <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, provide ID #: _____	Medicaid: <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, provide ID #: _____

C. PRESCRIBER INFORMATION

First Name:		Last Name: _____ (Check One): <input type="checkbox"/> M.D. <input type="checkbox"/> D.O. <input type="checkbox"/> N.P. <input type="checkbox"/> P.A.			
Address:		City:		State:	ZIP:
Phone:	Fax:	St Lic #:	NPI #:	DEA #:	UPIN:
Provider Email:		Office Contact Name:			Phone:
Specialty (Check one): <input type="checkbox"/> Oncologist <input type="checkbox"/> Hematologist <input type="checkbox"/> Other: _____					

D. DISPENSING PROVIDER/ADMINISTRATION INFORMATION

Place of Administration: <input type="checkbox"/> Self-administered <input type="checkbox"/> Physician's Office <input type="checkbox"/> Outpatient Infusion Center Phone: _____ Center Name: _____ <input type="checkbox"/> Home Infusion Center Phone: _____ Agency Name: _____ <input type="checkbox"/> Administration code(s) (CPT): _____ Address: _____	Dispensing Provider/Pharmacy: Patient Selected choice <input type="checkbox"/> Physician's Office <input type="checkbox"/> Retail Pharmacy <input type="checkbox"/> Specialty Pharmacy <input type="checkbox"/> Other Name: _____ Address: _____ Phone: _____ Fax: _____ TIN: _____ PIN: _____
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E. PRODUCT INFORMATION

Request is for: Enjaymo (sutimlimab-jome) Dose: _____ Frequency: _____

F. DIAGNOSIS INFORMATION - Please indicate primary ICD code and specify any other where applicable.

Primary ICD Code: _____ Secondary ICD Code: _____ Other ICD Code: _____

G. CLINICAL INFORMATION - Required clinical information must be completed in its entirety for all precertification requests.

For ALL Requests (clinical documentation required):

Yes No Is this infusion request in an outpatient hospital setting?

Yes No Has the patient experienced an adverse event with the requested product that has not responded to conventional interventions (e.g., acetaminophen, steroids, diphenhydramine, fluids, other pre-medications or slowing of infusion rate) or a severe adverse event (anaphylaxis, anaphylactoid reactions, myocardial infarction, thromboembolism, or seizures) during or immediately after an infusion?

Yes No Does the patient have severe venous access issues that require the use of special interventions only available in the outpatient hospital setting?

Yes No 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?
Please provide a description of the behavioral issue or impairment: _____

Yes No 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?
Please provide a description of the condition: Cardiopulmonary: _____
 Respiratory: _____
 Renal: _____
 Other: _____

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Patient First Name	Patient Last Name	Patient Phone	Patient DOB
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G. CLINICAL INFORMATION (continued) – Required clinical information must be completed in its entirety for all precertification requests.

For Initiation Requests (clinical documentation required for all requests):

- Yes No Was the diagnosis of primary cold agglutinin disease (CAD) confirmed by evidence of hemolysis?
- Yes No Does the patient have a lactate dehydrogenase (LDH) level above the upper limit of normal?
Please indicate level: _____ U/L
- Yes No Does the patient have a haptoglobin level below the lower limit of normal?
Please indicate level: _____ mg/L
- Yes No Does the patient have a positive polyspecific direct antiglobulin test (DAT) result?
- Yes No Does the patient have a monospecific direct antiglobulin test (DAT) result strongly positive for C3d?
- Yes No Does the patient have a cold agglutinin titer of 1:64 or higher measured at 4°C?
- Yes No Does the patient have a DAT IgG level of 1+ or less?
- Yes No 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)?

For Continuation Requests (clinical documentation required for all requests):

- Yes No Has patient experienced disease progression or unacceptable toxicity while on the current regimen?
- Yes No 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)?

H. ACKNOWLEDGEMENT

Request Completed By (Signature Required): _____ **Date:** ____/____/____

Any person who knowingly files a request for authorization of coverage of a medical procedure or service with the intent to injure, defraud or deceive any insurance company by providing materially false information or conceals material information for the purpose of misleading, commits a fraudulent insurance act, which is a crime and subjects such person to criminal and civil penalties.

The plan may request additional information or clarification, if needed, to evaluate requests.