



Uplizna™ (inebilizumab-cdon)
Medication Precertification Request

Page 1 of 2
 (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:
Phone: 1-866-503-0857
FAX: 1-844-268-7263

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:	<i>(Check One):</i> <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"/> Neurologist <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 Uplizna (inebilizumab-cdon) 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 for all requests):

Yes No Does the patient have a diagnosis of neuromyelitis optica spectrum disorder (NMOSD)?

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?

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?

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

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

Yes No Is the patient anti-aquaporin-4 (AQP4) antibody positive?

Yes No Will the requested drug be used concomitantly with rituximab or eculizumab (Soliris)?

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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.

Please indicate which of the following applies to the patient:

- Optic neuritis
- Acute myelitis
- Area postrema syndrome (episode of otherwise unexplained hiccups or nausea and vomiting)
- Acute brainstem syndrome
- Symptomatic narcolepsy or acute diencephalic clinical syndrome with NMOSD-typical diencephalic magnetic resonance imaging (MRI) lesions
- Symptomatic cerebral syndrome with NMOSD-typical brain lesions
- None of the above

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

- Yes No Has the patient demonstrated a positive response to therapy (e.g., reduction in number of relapses)?

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.