Rituxan Hycela® (rituximab and hyaluronidase human) Medication Precertification Request

(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 #: _____
Group #: _____
Insured:

Does patient have other coverage? ☐ Yes ☐ No
If yes, provide ID#: _____ Carrier Name: _____
Insured:

Medicare: ☐ Yes ☐ No If yes, provide ID #:

Medicaid: ☐ Yes ☐ No If yes, provide ID #:

C. PRESCRIBER INFORMATION

First Name:		Last Name:		*(Check One):* ☐ M.D. ☐ D.O. ☐ N.P. ☐ P.A.	
Address:			City:	State:	ZIP:
Phone:	Fax:	St Lic #:	NPI #:	DEA #:	UPIN:
Provider Email:		Office Contact Name:		Phone:	

Specialty *(Check one):* ☐ **Oncologist** ☐ **Other**: _____

D. DISPENSING PROVIDER/ADMINISTRATION INFORMATION

Place of Administration:

☐ Self-administered ☐ Physician's Office
☐ Outpatient Infusion Center Phone: _____
Center Name: _____
☐ Home Infusion Center Phone: _____
Agency Name: _____
☐ Administration code(s) (CPT): _____
Address: _____

Dispensing Provider/Pharmacy: *Patient Selected choice*

☐ Physician's Office ☐ Retail Pharmacy
☐ Specialty Pharmacy ☐ Other _____
Name: _____
Address: _____
Phone: _____ Fax: _____
TIN: _____ PIN: _____

E. PRODUCT INFORMATION

Request is for: Rituxan Hycela (rituximab and hyaluronidase) **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 Has the patient failed treatment with Truxima due to a documented intolerable adverse event (e.g., rash, nausea, vomiting)?
→ ☐ Yes ☐ No Was the adverse event unexpected and not attributed to the active ingredient as described in the prescribing information (i.e., known adverse reaction for both the brand and biosimilar medication)?

Please indicate the patient's documented diagnosis:

☐ Castleman's disease (CD)
☐ Chronic lymphocytic leukemia (CLL)
☐ Diffuse large B-cell lymphoma (DLBCL)
☐ Follicular lymphoma (FL)
☐ Gastric mucosa-associated lymphoid tissue (MALT) lymphoma
☐ Hairy cell leukemia
☐ High-grade B-cell lymphoma
☐ Histologic transformation of indolent lymphomas to diffuse large B-cell lymphoma
☐ Mantle cell lymphoma
☐ Nodal marginal zone lymphoma
☐ Nongastric MALT lymphoma
☐ Post-transplant lymphoproliferative disorder (PTLD)
☐ Primary cutaneous B-cell lymphoma (e.g., cutaneous marginal zone lymphoma or cutaneous follicle center lymphomas)
☐ Small lymphocytic lymphoma (SLL)
☐ Splenic marginal zone lymphoma

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Patient First Name	Patient Last Name	Patient Phone	Patient DOB

G. CLINICAL INFORMATION *(continued)* – Required clinical information must be completed in its entirety for all precertification requests.

For Initiation Requests (clinical documentation required):

☐ Yes ☐ No Does the patient have CD20 positive disease that was confirmed by testing or analysis?

☐ Yes ☐ No Has the patient received at least one full dose of a rituximab product by IV infusion without experiencing severe adverse reactions?

For Continuation Requests (clinical documentation required):

☐ Yes ☐ No Is there evidence of unacceptable toxicity while on the current regimen?

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.