Kyprolis® (carfilzomib) Medication Precertification Request

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

Aetna Precertification Notification
Phone: 1-866-752-7021 (TTY: 711)
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: ☐ **Kyprolis (carfilzomib) 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 Multiple Myeloma Requests (clinical documentation required for all requests):
Please indicate the patient's Body Surface Area (BSA): _____m^2
For once weekly treatment:
☐ Yes ☐ No Will the patient's dose exceed 70 mg/m2 (not to exceed 154 mg per dose)?
☐ Yes ☐ No Will the patient be receiving more than 3 doses per 28 days?
For twice weekly treatment:
☐ Yes ☐ No Will the patient's dose exceed 56 mg/m2 (not to exceed 124 mg per dose)?
☐ Yes ☐ No Will the patient be receiving more than 6 doses per 28 days?

For Initiation Requests (clinical documentation required for all requests):
☐ **Multiple myeloma**
☐ Yes ☐ No Has the patient had a contraindication, intolerance or ineffective response to Velcade or its generic equivalent bortezomib?
Please indicate the prescribed regimen:
☐ The requested medication in combination with dexamethasone
→ What is the clinical setting in which the requested medication will be used? ☐ Progressive disease ☐ Refractory disease ☐ Relapsed disease ☐ Other _____
☐ The requested medication in combination with cyclophosphamide and dexamethasone
☐ The requested medication in combination with lenalidomide and dexamethasone
☐ The requested medication in combination with daratumumab, lenalidomide and dexamethasone
☐ The requested medication in combination with daratumumab and dexamethasone
→ What is the clinical setting in which the requested medication will be used? ☐ Progressive disease ☐ Refractory disease ☐ Relapsed disease ☐ Other _____

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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.

☐ The requested medication in combination with daratumumab, hyaluronidase-fihj and dexamethasone
→ What is the clinical setting in which the requested medication will be used? ☐ Progressive disease ☐ Refractory disease ☐ Relapsed disease ☐ Other _____

☐ The requested medication in combination with pomalidomide and dexamethasone
→ What is the clinical setting in which the requested medication will be used? ☐ Relapsed disease ☐ Progressive disease ☐ Other

☐ The requested medication in combination with cyclophosphamide, thalidomide, and dexamethasone
→ What is the clinical setting in which the requested medication will be used? ☐ Relapsed disease ☐ Progressive disease ☐ Other

☐ The requested medication in combination with isatuximab-irfc and dexamethasone
→ What is the clinical setting in which the requested medication will be used? ☐ Progressive disease ☐ Refractory disease ☐ Relapsed disease ☐ Other _____

☐ The requested medication in combination with selinexor and dexamethasone
→ What is the clinical setting in which the requested medication will be used? ☐ Progressive disease ☐ Relapsed disease ☐ Other

☐ The requested medication will be used as a single agent
→ ☐ Yes ☐ No Has the patient received at least one prior therapy?

☐ The requested medication in combination with lenalidomide
→ ☐ Yes ☐ No Will the requested medication be used as a maintenance therapy for symptomatic disease?

☐ The requested medication in combination with bendamustine and dexamethasone
→ ☐ Yes ☐ No Has the patient received more than 3 prior therapies?
What is the clinical setting in which the requested medication will be used? ☐ Progressive disease ☐ Relapsed disease ☐ Other

☐ Other _____

☐ **Systemic light chain amyloidosis**
☐ Yes ☐ No What is the clinical setting in which the requested medication will be used? ☐ Relapsed disease ☐ Refractory disease ☐ Other

☐ **Waldenstrom macroglobulinemia/lymphoplasmacytic lymphoma**

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

☐ Yes ☐ No Has the patient experienced unacceptable toxicity or disease progression 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.