



MEDICARE FORM

Kyprolis (carfilzomib) Medication Precertification Request

Page 1 of 2

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

Virginia (HMO D-SNP)
FAX: 1-833-280-5224
PHONE: 1-855-463-0933

For other lines of business:
Please use other form.

**Note: Kyprolis is non-preferred.
Bortezomib and Velcade are preferred.**

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"/> Other: _____					

D. DISPENSING PROVIDER/ADMINISTRATION INFORMATION

Place of Administration:		Dispensing Provider/Pharmacy: Patient Selected choice	
<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: _____		<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: _____	

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): ____m²
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):
Note: Kyprolis is non-preferred. Bortezomib and Velcade are preferred.
 Yes No Has the patient had prior therapy with Kyprolis within the last 365 days?
 Yes No Has the patient had a trial and failure, intolerance, or contraindication to any of the following? (select all that apply)
 Bortezomib Velcade (bortezomib)
Please explain if there are any other medical reason(s) that the patient cannot use any of the following preferred products when indicated for the patient's diagnosis? (select all that apply)
 Bortezomib Velcade (bortezomib)

Continued on next page



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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.

Multiple myeloma

Please indicate the prescribed regimen:

- The requested medication in combination with dexamethasone
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
The requested medication in combination with daratumumab and hyaluronidase-fihj and dexamethasone
The requested medication in combination with panobinostat
The requested medication in combination with pomalidomide and dexamethasone
The requested medication in combination with cyclophosphamide, thalidomide, and dexamethasone
The requested medication in combination with isatuximab-irfc and dexamethasone
The requested medication in combination with selinexor and dexamethasone
The requested medication as a single agent

- Systemic light chain amyloidosis
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.