

Kyprolis

CareFirst Prior Authorization Request

CVS Caremark administers the prescription benefit plan for the patient identified. This patient's benefit plan requires prior authorization for certain medications in order for the drug to be covered. To make an appropriate determination, providing the most accurate diagnosis for the use of the prescribed medication is necessary. **Please respond below and fax this form to CVS Caremark toll-free at 1-855-330-1720**. If you have questions regarding the prior authorization, please contact CVS Caremark at **1-888-877-0518**. For inquiries or questions related to the patient's eligibility, drug copay or medication delivery; please contact the Specialty Customer Care Team: CaremarkConnect® 1-800-237-2767.

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Patient's Name:	Date:
Patient's ID:	
Specialty:	
Physician Office Telephone:	
<u>Referring</u> Provider Info: 🛛 Same as Reque Name:	0
Fax:	
	ring Provider 🖵 Same as Requesting Provider
Name:	NPI#:
Fax:	Phone:

Approvals may be subject to dosing limits in accordance with FDA-approved labeling, accepted compendia, and/or evidence-based practice guidelines.

Required Demographic Information:

 Patient Weight:
 kg

 Patient Height:
 cm

Please indicate the place of service for the requested drug: Ambulatory Surgical Home On Campus Outpatient Hospital Office

Off Campus Outpatient Hospital
 Pharmacy

What is the ICD-10 code?

Send completed form to: Case Review Unit CVS Caremark Specialty Programs Fax: 1-855-330-1720

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CVS Caremark Specialty Pharmacy • 2211 Sanders Road NBT-6 • Northbrook, IL 60062

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Criteria Questions:

1. What is the diagnosis?

□ Multiple myeloma, *Continue to 2*

Systemic light chain amyloidosis, *Continue to 2*

Waldenstrom macroglobulinemia/lymphoplasmacytic lymphoma, *Continue to 2* POEMS (polyneuropathy, organomegaly, endocrinopathy, monoclonal protein, skin changes) syndrome,

Continue to 2

□ Other, please specify. _____, Continue to 2

2. Is this a request for continuation of therapy with the requested medication?

□ Yes, *Continue to 3*

 \square No, Continue to 4

3. Has the patient experienced unacceptable toxicity or disease progression while on the current regimen? Yes, *Continue to 22*

■ No, *Continue to 22*

4. What is the diagnosis?

□ Multiple myeloma, *Continue to 5*

□ Systemic light chain amyloidosis, *Continue to 22*

□ Waldenstrom macroglobulinemia/lymphoplasmacytic lymphoma, *Continue to 22*

Dependence of POEMS (polyneuropathy, organomegaly, endocrinopathy, monoclonal protein, skin changes) syndrome, *Continue to 21*

5. What is the prescribed regimen?

 \Box The requested medication in combination with dexamethasone, *Continue to 6*

The requested medication in combination with cyclophosphamide and dexamethasone, Continue to 22

The requested medication in combination with lenalidomide and dexamethasone, Continue to 22

The requested medication in combination with daratumumab, lenalidomide and dexamethasone, *Continue to* 22

The requested medication in combination with daratumumab and dexamethasone, Continue to 7

 \Box The requested medication in combination with daratumumab, hyaluronidase-fihj and dexamethasone, *Continue* to 8

The requested medication in combination with pomalidomide and dexamethasone, Continue to 9

 \Box The requested medication in combination with pomalidomide, daratumumab, and dexamethasone, *Continue to* 9

□ The requested medication in combination with cyclophosphamide, thalidomide, and dexamethasone, *Continue to 10*

The requested medication in combination with isatuximab-irfc and dexamethasone, *Continue to 11*

The requested medication in combination with selinexor and dexamethasone, Continue to 12

□ The requested medication as a single agent, *Continue to 13*

The requested medication in combination with lenalidomide, Continue to 14

The requested medication in combination with bendamustine and dexamethasone, Continue to 15

The requested medication in combination with venetoclax and dexamethasone, Continue to 17

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□ The requested medication in combination with isatuximab-irfc, lenalidomde, and dexamethasone, *Continue to* 19

□ Other, please specify. , *Continue to 22* 6. What is the clinical setting in which the requested medication will be used? □ Relapsed disease, *Continue to 22* **□** Refractory disease, *Continue to 22* □ Progressive disease, *Continue to 22* □ Other, please specify. _____, *Continue to 22* 7. What is the clinical setting in which the requested medication will be used? □ Relapsed disease, *Continue to 22* **□** Refractory disease, *Continue to 22* □ Progressive disease, *Continue to 22* □ Other, please specify. _____, *Continue to 22* 8. What is the clinical setting in which the requested medication will be used? **Relapsed disease**, *Continue to 22* **Refractory disease**, *Continue to 22* □ Progressive disease, *Continue to 22* □ Other, please specify. _____, *Continue to 22* 9. What is the clinical setting in which the requested medication will be used? **Relapsed disease**, *Continue to 22* □ Progressive disease, *Continue to 22* □ Other, please specify. , *Continue to 22* 10. What is the clinical setting in which the requested medication will be used? **Relapsed disease**. *Continue to 22* □ Progressive disease, *Continue to 22* □ Other, please specify. _____, *Continue to 22* 11. What is the clinical setting in which the requested medication will be used? **Relapsed disease**, *Continue to 22* **Refractory disease**, *Continue to 22* □ Progressive disease, *Continue to 22* □ Other, please specify. _____, *Continue to 22* 12. What is the clinical setting in which the requested medication will be used?

□ Relapsed disease, *Continue to 22*

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□ Progressive disease, *Continue to 22* □ Other, please specify. _____, Continue to 22 13. Has the patient received at least one prior therapy? □ Yes, Continue to 22 □ No, *Continue to 22* 14. Will the requested medication be used as maintenance therapy for symptomatic disease? □ Yes, Continue to 22 \square No, *Continue to 22* 15. Has the patient received more than 3 prior therapies? □ Yes, Continue to 16 □ No, Continue to 16 16. What is the clinical setting in which the requested medication will be used? □ Relapsed disease, *Continue to 22* □ Refractory disease, *Continue to 22* □ Other, please specify. _____, Continue to 22 17. What is the clinical setting in which the requested medication will be used? □ Relapsed disease, *Continue to 18* □ Progressive disease, *Continue to 18* □ Other, please specify. _____, Continue to 18 18. Does the patient have a documented t(11:14) translocation? ACTION REQUIRED: If Yes, attach chart note(s) or test results of t(11:14) translocation. □ Yes ACTION REQUIRED: Submit supporting documentation, Continue to 22 \square No, *Continue to 22* **U**nknown, *Continue to 22* 19. Will the requested medication be used as primary therapy for symptomatic disease? \Box Yes. Continue to 20 \square No, Continue to 20 20. Is the patient a candidate for transplant? □ Yes, Continue to 22 \square No, *Continue to 22* 21. Will the requested medication be used in combination with dexamethasone? \Box Yes, Continue to 22 □ No, Continue to 22

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22. What is the patient's height in inches? ______ inches, Continue to 23

23. What is the patient's weight in pounds? pounds, *Continue to 24*

24. What is the patient's Body Surface Area (BSA) (Note: average adult BSA is around 1.7 m2)? _____m2, Continue to 25

25. What is the patient's dose in milligrams? _____ mg, Continue to 26

26. How frequently will the patient be receiving the requested medication?

□ Once weekly, *Continue to 27*

Twice weekly, Continue to 29

27. Will the patient's dose exceed 70 mg/m2 (not to exceed 154 mg per dose)?

□ Yes, *Continue to 28*

 \square No, Continue to 28

28. Will the patient be receiving more than 3 doses per 28 days?

□ Yes, No Further Questions

□ No, No Further Questions

29. Will the patient's dose exceed 56 mg/m2 (not to exceed 124 mg per dose)?

□ Yes, Continue to 30

□ No. Continue to 30

30. Will the patient be receiving more than 6 doses per 28 days? □ Yes, No Further Questions **D** No, No Further Questions

I attest that this information is accurate and true, and that documentation supporting this information is available for review if requested by CVS Caremark or the benefit plan sponsor.

Х

Prescriber or Authorized Signature

Date (mm/dd/yy)

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