



Proleukin (aldesleukin)

Prior Authorization Request

Your patient's benefit plan requires prior authorization for certain medications. In order to make appropriate medical necessity determinations, your patient's diagnosis and other clinical information is required. **Please complete the information requested on the form below and fax this form to Priority Partners, toll-free at 1-866-212-4756** to initiate the review process. If you have questions regarding the prior authorization please contact Priority Partners at 888-819-1043 Option 4.

Patient's Name: _____ Date: _____
Patient's ID: _____ Patient's Date of Birth: _____
Physician's Name: _____
Specialty: _____ NPI#: _____
Physician Office Telephone: _____ Physician Office Fax: _____

Referring Provider Info: Same as Requesting Provider

Name: _____ NPI#: _____
Fax: _____ Phone: _____

Rendering Provider Info: Same as Referring 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 Off Campus Outpatient Hospital
 On Campus Outpatient Hospital Office Pharmacy

Drug Information:

Strength/Measure _____ Units ml Gm mg ea Un

Directions(sig) _____ Route of administration _____

Dosing frequency _____

Send completed form to: Priority Partners Fax: 1-866-212-4756

Note: This fax may contain medical information that is privileged and confidential and is solely for the use of individuals named above. If you are not the intended recipient you hereby are advised that any dissemination, distribution, or copying of this communication is prohibited. If you have received the fax in error, please immediately notify the sender by telephone and destroy the original fax message. Proleukin [aldesleukin] SGM – 12/2022.

Priority Partners • 7231 Parkway Drive Suite 100 • Hanover, MD 21076

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Criteria Questions:

1. What is the diagnosis?

- Renal cell carcinoma (If checked, go to 2)
- Cutaneous melanoma (If checked, go to 2)
- Chronic graft-versus-host disease (GVHD) (If checked, go to 2)
- Neuroblastoma (If checked, go to 2)
- Other, please specify. _____ (If checked, go to 2)

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

- Yes, *Continue to 3*
- No, *Continue to 10*

3. What is the diagnosis?

- Renal cell carcinoma (If checked, go to 4)
- Cutaneous melanoma (If checked, go to 4)
- Chronic graft-versus-host disease (GVHD) (If checked, go to 8)
- Neuroblastoma (If checked, go to 9)

4. Has the patient been evaluated for response approximately 4 weeks after completion of a course of therapy with the requested drug and will again be evaluated immediately prior to the scheduled start of the next treatment course?

- Yes, *Continue to 5*
- No, *Continue to 5*

5. Did the patient experience any tumor shrinkage following the last course of therapy with the requested drug?

- Yes, *Continue to 6*
- No, *Continue to 6*

6. Is retreatment with the requested drug contraindicated for the patient?

- Yes, *Continue to 7*
- No, *Continue to 7*

7. Will the patient's treatment course with the requested drug be separated by a rest period of at least 7 weeks from the date of hospital discharge?

- Yes, *No Further Questions*
- No, *No Further Questions*

8. Is there improvement in symptoms and no unacceptable toxicity?

- Yes, *No Further Questions*
- No, *No Further Questions*

9. Is there evidence of unacceptable toxicity or disease progression while on the current regimen?

- Yes, *No Further Questions*
- No, *No Further Questions*

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10. What is the diagnosis?

- Renal cell carcinoma (If checked, go to 11)
- Cutaneous melanoma (If checked, go to 12)
- Chronic graft-versus-host disease (If checked, go to 14)
- Neuroblastoma (If checked, *no further questions*)

11. What is the clinical setting in which the requested drug will be used?

- Metastatic disease (If checked, *no further questions*)
- Other, please specify. _____ (If checked, *no further questions*)

12. What is the clinical setting in which the requested drug will be used?

- Metastatic disease (If checked, go to 13)
- Unresectable disease (If checked, go to 13)
- Other, please specify. _____ (If checked, go to 13)

13. Will the requested drug be given as high-dose single agent therapy for subsequent therapy?

- Yes, *No Further Questions*
- No, *No Further Questions*

14. Did the patient respond to first-line therapy options?

- Yes (If checked, go to 15)
- No (If checked, go to 15)
- Unknown (If checked, go to 15)

15. Is the requested drug being used as additional therapy in conjunction with systemic corticosteroids?

- Yes, *No Further Questions*
- 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.

X

Prescriber or Authorized Signature

Date (mm/dd/yy)

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