

Targretin (bexarotene)

Override(s)	Approval Duration
Prior Authorization Quantity Limit	1 year

Medications	Quantity Limit
Targretin (bexarotene) 75mg capsules	May be subject to quantity limit
Targretin (bexarotene) 1% gel	

APPROVAL CRITERIA

Requests for oral Targretin (bexarotene) may be approved if the following criteria are met:

I. Individual is under 19 years of age;

OR

II. Individual has a diagnosis of one of the following:

A. Mycosis Fungoides/Sézary syndrome (NCCN 2A);

OR

B. Primary cutaneous CD30+ T-cell lymphoproliferative disorder (for example, primary cutaneous anaplastic large-cell lymphoma/ALCL, lymphomatoid papulosis/LyP) (NCCN 2A); **AND**

1. Individual is using for primary treatment; **OR**

2. Individual is using in relapsed/refractory disease;

OR

C. Other cutaneous T-Cell lymphomas where disease is refractory to one other prior non-topical therapy;

OR

D. Subcutaneous panniculitis-like T-cell lymphoma and individual is using in one of the following regimens (NCCN 2A):

1. As a single agent or in combination with prednisone in individuals without hemophagocytic lymphohistiocytosis and with low tumor burden;

OR

2. As maintenance therapy as a single agent following complete or partial response to first line therapy.

Requests for Targretin (bexarotene) 1% gel may be approved if the following criteria are met:

- I. Individual is under 19 years of age;

OR

- II. Individual has a diagnosis of one of the following:

- A. Cutaneous T-cell leukemia/lymphoma; **AND**

1. Individual is using in the topical treatment of cutaneous lesions; **AND**
 2. Individual has refractory or persistent disease after other therapies;
- OR**
3. Individual has intolerance to other therapies;

OR

- B. Primary cutaneous B-cell lymphoma; **AND**

1. Individual has primary cutaneous marginal zone or follicle center lymphoma (NCCN 2A);

OR

- C. Mycosis Fungoides/Sézary syndrome (NCCN 2A);

OR

- D. Adult-T Cell Leukemia/Lymphoma; **AND**

1. Individual is using for chronic/smoldering subtype as first-line skin-directed therapy (NCCN 2A).

Key References:

1. DailyMed. Package inserts. U.S. National Library of Medicine, National Institutes of Health website. <http://dailymed.nlm.nih.gov/dailymed/about.cfm>. Updated periodically.
2. DrugPoints® System [electronic version]. Truven Health Analytics, Greenwood Village, CO. Updated periodically.
3. Lexi-Comp ONLINE™ with AHFS™, Hudson, Ohio: Lexi-Comp, Inc.; 2025; Updated periodically.
4. NCCN Clinical Practice Guidelines in Oncology™. © 2025 National Comprehensive Cancer Network, Inc. For additional information visit the NCCN website: <http://www.nccn.org/index.asp>. Accessed on October 9, 2025.
 - a. Primary Cutaneous Lymphomas. V3.2025. Revised June 25, 2025.
 - b. T-Cell Lymphomas. V2.2025. Revised May 28, 2025.

Federal and state laws or requirements, contract language, and Plan utilization management programs or policies may take precedence over the application of this clinical criteria.

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