Introduction

The lymphatic system, which is part of the immune system, is composed of specialized cells. Most of these specialized cells are white blood cells known as lymphocytes. There are two main types of lymphocytes: B-cells and T-cells. Lymphoma is cancer that starts in lymphocytes.

Cutaneous T-cell lymphomas are cancers that start in the T-cell lymphocytes and affect the skin. Mycosis fungoides and Sézary syndrome are two examples of lymphomas of the skin.

T-cell lymphoma can arise in other parts of the body such as lymph nodes, lining of the intestines, or the spleen, liver, or colon.

Treating T-cell lymphomas depends on the specific type of cancer and how slow or fast it’s growing. This policy describes when particular chemotherapy drugs may be considered medically necessary for T-cell lymphomas that did not respond to initial treatment.

Note: The Introduction section is for your general knowledge and is not to be taken as policy coverage criteria. The rest of the policy uses specific words and concepts familiar to medical professionals. It is intended for providers. A provider can be a person, such as a doctor, nurse, psychologist, or dentist. A provider also can be a place where medical care is given, like a hospital, clinic, or lab. This policy informs them about when a service may be covered.

Policy Coverage Criteria
**Note:** Initial approval period for agents listed below will be 3 months. Continued approval beyond the first 3 months will require documentation showing objective response to therapy.

<table>
<thead>
<tr>
<th>Drug</th>
<th>Medical Necessity</th>
</tr>
</thead>
</table>
| Targretin® (bexarotene), Zolinza® (vorinostat) or Istodax® (romidepsin) | Targretin® (bexarotene), Zolinza® (vorinostat), or Istodax® (romidepsin) may be considered medically necessary in patients who are refractory to at least one prior systemic therapy and have:  
  - Cutaneous or peripheral T-cell lymphoma
  OR  
  - Mycosis fungoides/Sezary syndrome  
  
  **AND**  
  Systemic therapies (including but not limited to the following list of agents recommended by NCCN guidelines):  
  - For simple skin involvement:  
    o Systemic retinoids (isotretinoin, acitretin, etc.)  
    o Interferons (alpha, gamma)  
    o Methotrexate  
    o Campath® (alemtuzumab)  
  OR  
  - For late stage disease usually with solid organ involvement, more aggressive therapy is usually required, for example:  
    o Gemcitabine  
    o Liposomal doxorubicin  

  Targretin® (bexarotene) topical gel may be approved when at least 3 of the following therapies have failed (including but not limited to the following list of agents recommended by NCCN guidelines):  
  - Phototherapy (UVB, NB-UVB, PUVA)  
  - Topical imiquimod  
  - Topical corticosteroids  
  - Topical mechlorethamine  
  - Local radiation  

| Beleodaq® (belinostat) | Beleodaq® (belinostat) may be considered medically necessary for the treatment of patients with relapsed or refractory |
Drug | Medical Necessity
---|---
angioimmunoblastic T-cell lymphoma, anaplastic large cell lymphoma, enteropathy-associated T-cell lymphoma or other peripheral T-cell lymphoma who have failed or are refractory to at least one prior systemic therapy.

**Valchlor® (mechlorethamine)**

Valchlor® (mechlorethamine) may be considered medically necessary for topical treatment of Stage IA and IB mycosis fungoides-type cutaneous T-cell lymphoma in patients who have tried and failed at least 2 of the following therapies (as recommended by NCCN guidelines):
- Topical corticosteroids (prednisone, triamcinolone, etc.)
- Topical imiquimod
- Phototherapy
- Local radiation

All other uses of the medications listed in this policy are considered investigational.

### Coding

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>HCPCS</strong></td>
<td></td>
</tr>
<tr>
<td>J9032</td>
<td>Injection, belinostat (Beleodaq®), 10 mg</td>
</tr>
<tr>
<td>J9315</td>
<td>Injection, romidepsin (Istodax®), 1 mg</td>
</tr>
</tbody>
</table>

**Note:** CPT codes, descriptions and materials are copyrighted by the American Medical Association (AMA). HCPCS codes, descriptions and materials are copyrighted by Centers for Medicare Services (CMS).

### Related Information

### Benefit Application

The drugs included in this policy may be covered under either the pharmacy or medical benefit.
**Evidence Review**

**Description**

**Cutaneous T-Cell Lymphoma**

Cutaneous T-cell lymphoma (CTCL) is a rare type of cancer of unknown etiology. In the United States, the estimated annual incidence of CTCL is 0.4 to 1 case per 100,000 (or 1,000 to 3,000 new cases per year). Since most cases are not fatal and patients can live with CTCL for 10 or more years, estimated U.S. prevalence is approximately 20,000. Approximately 25% of these cases (5,000) have advanced disease.

CTCL is characterized by abnormal malignant T-cells residing in the skin which cause skin lesions and pruritis. The disease often remains confined to the skin and has an indolent course. However, in more aggressive CTCL, the disease may progress to involve other areas of the body such as lymph nodes, blood, and organs. The most common indolent form is mycosis fungoides. This form usually manifests as flat red or pink scaly patches that first appear in sun-protected areas of the body (e.g., buttocks or trunk). As the disease progresses, the patches evolve into a raised pruritic plaque phase. Plaques can also evolve into nodular skin tumors which may ulcerate.

The most common aggressive form of CTCL is Sézary syndrome. This syndrome accounts for about 5% of all CTCL cases and the prognosis is poor. Median survival is usually less than 3 years. It is characterized by widespread erythroderma and the presence of abnormal CD4+ T-cells (called Sézary cells) in both the skin and peripheral blood. This condition may also be accompanied by scaling pruritic plaques or tumors. In advanced stages, patients with Sézary syndrome may develop alopecia, leonine facies, hyperkeratosis of the palms and soles, nail dystrophy, and parasthesia.

Therapeutic management of CTCL is based largely on the stage of the disease. In early, mild, and slowly progressive stages of the disease, treatment is mainly targeted at reducing symptoms and lesions with topical therapies (topical corticosteroids, topical retinoids (bexarotene, tazarotene), topical chemotherapy (nitrogen mustard, carmustine), local radiation (for limited skin involvement) or electron beam therapy (for severe skin involvement), phototherapy (UVB, nbUVB or patch/thin plaques; PUVA for thicker plaques), topical imiquimod (for limited localized skin involvement). In patients with advanced stages of the disease, mycosis fungoides and Sézary syndrome, systemic therapies, such as retinoids (e.g., bexarotene, acitretin), may be employed.
Targretin® (bexarotene) is a member of a subclass of retinoids that selectively activate retinoid X receptors (RXRs). These retinoid receptors have biologic activity distinct from that of retinoic acid receptors (RARs). Retinoids are associated with birth defects in humans. Targretin® (bexarotene) also caused birth defects when administered orally to pregnant rats. Therefore, it must not be administered during pregnancy.

Targretin® (bexarotene) selectively binds and activates retinoid X-receptor subtypes (RXRα, RXRβ, RXRγ). RXRs can form heterodimers with various receptor partners such as retinoic acid receptors (RARs), vitamin D receptor, thyroid receptor, and peroxisome proliferator activator receptors (PPARs). Once activated, these receptors function as transcription factors that regulate the expression of genes that control cellular differentiation and proliferation. Targretin® (bexarotene) inhibits the growth in vitro of some tumor cell lines of hematopoietic and squamous cell origin. It also induces tumor regression in vivo in some animal models. The exact mechanism of action of Targretin® (bexarotene) in the treatment of cutaneous T-cell lymphoma (CTCL) is unknown.

Bexarotene (Targretin) is indicated for the treatment of cutaneous manifestations of cutaneous T cell lymphoma in patients who are refractory to at least one prior systemic therapy.

Zolinza® (vorinostat) and Istodax® (romidepsin) are a histone deacetylase (HDAC) inhibitors with enzymatic inhibitory activity at nanomolar concentrations for HDAC1, HDAC2, HDAC3 [class I], and HDAC6 [class II]. These enzymes catalyze the removal of acetyl groups from the lysine residues of proteins, including histones and transcription factors. In some cancer cells, HDACs are overexpressed or are aberrantly recruited for oncogenic transcription causing hypoacetylation of core nucleosomal histones. While the exact mechanism of action of these agents has not been fully characterized, in vitro, they cause accumulation of acetylated histones, induces cell cycle arrest, and/or apoptosis of some transformed cells.

Zolinza® (vorinostat) is indicated for treatment of cutaneous manifestations in patients with cutaneous T-cell lymphoma (CTCL) who have progressive, persistent or recurrent disease on or following two systemic therapies.

Istodax® (romidepsin) is indicated for treatment of cutaneous T-cell lymphoma (CTCL) or peripheral T-cell lymphoma in patients who have received at least one prior systemic therapy.
Rationale

*Targretin® (bexarotene)*

**Efficacy**

No randomized controlled trial for *Targretin® (bexarotene)* in CTCL was found on literature search. The efficacy of oral bexarotene was demonstrated in 2 pivotal multinational, open-label, phase II-III studies in 94 adults with advanced refractory CTCL and 58 with early stage refractory or persistent CTCL. At recommended dosing (300 mg/m2/day), overall response to monotherapy as measured by PGA and CAILDS was about 50%. Two supportive open-label studies show systemic combination therapy to be efficacious as measured by response rates.

The efficacy of topical bexarotene was demonstrated in 1 pivotal multinational, open-label, phase II-III study in 50 adults with treatment refractory early stage CTCL. Like systemic bexarotene, overall response was about 50%. Bexarotene gel is the only topical product treatment of CTCL which has received approval by the U.S. Food and Drug Administration (FDA).

Off-label studies for systemic bexarotene have been reported for prevention of breast cancer, head and neck cancer, psoriasis, non-small cell lung cancer, and acute myeloid leukemia. Off-label studies for topical bexarotene have been reported for alopecia areata, parapsoriasis, severe hand dermatitis, and psoriasis.

No comparative studies between bexarotene and therapeutic alternatives for CTCL have been published and no comparative trials are currently underway.

**Safety**

*Targretin® (bexarotene)* carries a boxed warning the drug should not be used in pregnant females because of its association with birth defects. Systemic bexarotene induces major lipid abnormalities in most patients, particularly hypertriglyceridemia. Lipids should be monitored at baseline, weekly for 2-4 weeks, then at 8 week intervals. Abnormalities are reversible with cessation of therapy and can be managed with dose reduction or concomitant antilipemic agents. Other serious adverse events reported following use of systemic bexarotene are pancreatitis, liver function test abnormalities and hepatotoxicities, hypothyroidism, leukopenia, and cataracts. Regular monitoring of related laboratory measures and ophthalmologic evaluations are recommended.
In oral CTCL trials, adverse events leading to dose reduction or study drug discontinuation in at least two patients were hyperlipemia, neutropenia/leukopenia, diarrhea, fatigue/lethargy, hypothyroidism, headache, liver function test abnormalities, rash, pancreatitis, nausea, anemia, allergic reaction, muscle spasm, pneumonia, and confusion. In topical CTCL trials, adverse events leading to dose reduction or study drug discontinuation in at least two patients were rash, contact dermatitis, and pruritus.

In 504 patients participating in oral CTCL and non-CTCL clinical trials, 3 patients experienced a severe adverse event that was fatal; one each from acute pancreatitis, a subdural hematoma, and liver failure. In topical CTCL trials, only one patient (2%) experienced a severe adverse event (rash).

The most common adverse events observed in patients with CTCL (n=86) receiving Zolinza® (vorinostat) 400 mg once-daily in uncontrolled clinical were diarrhea, fatigue, nausea, thrombocytopenia, anorexia, and dysgeusia. The most commonly reported serious adverse events, regardless of causality, were pulmonary embolism and anemia. In addition, laboratory abnormalities (eg, increased serum glucose, increased serum creatinine, proteinuria) were observed in all patients. Approximately 9% of the 86 patients discontinued the drug due to adverse events.

**Istodax® (romidepsin)**

**Efficacy**

Istodax® (romidepsin) was evaluated in 2 multicenter, single-arm clinical studies in patients with CTCL. Overall, 167 patients with CTCL were treated in the US, Europe, and Australia. Study 1 included 96 patients with confirmed CTCL after failure of at least 1 prior systemic therapy. Study 2 included 71 patients with a primary diagnosis of CTCL who received at least 2 prior skin directed therapies or one or more systemic therapies. Patients were treated with romidepsin at a starting dose of 14 mg/m² infused over 4 hours on days 1, 8, and 15 every 28 days. In both studies, patients could be treated until disease progression at the discretion of the investigator and local regulators. Objective disease response was evaluated according to a composite endpoint that included assessments of skin involvement, lymph node and visceral involvement, and abnormal circulating T-cells (“Sézary cells”). The primary efficacy endpoint for both studies was overall objective disease response rate (ORR) based on the investigator assessments, and defined as the proportion of patients with confirmed complete response (CR) or partial response (PR). CR was defined as no evidence of disease and PR as ≥50% improvement in disease. Secondary endpoints in both studies included duration of response and time to response.
Median time to first response was 2 months (range 1 to 6) in both studies. Median time to CR was 4 months in Study 1 and 6 months in Study 2 (range 2 to 9).

Istodax® (romidepsin) was evaluated in a multicenter, single-arm, international clinical study in patients with PTCL who had failed at least 1 prior systemic therapy (Study 3). Patients in US, Europe and Australia were treated with romidepsin at a dose of 14 mg/m2 infused over 4 hours on days 1, 8, and 15 every 28 days. Of the 131 patients treated, 130 patients had histological confirmation by independent central review and were evaluable for efficacy (HC Population). Six cycles of treatment were planned; patients who developed progressive disease (PD), significant toxicity, or who met another criterion for study termination were to discontinue treatment. Responding patients had the option of continuing treatment beyond 6 cycles at the discretion of the patient and Investigator until study withdrawal criteria were met. Primary assessment of efficacy was based on rate of complete response (CR + CRu) as determined by an Independent Review Committee (IRC) using the International Workshop Response Criteria (IWC). Secondary measures of efficacy included IRC assessment of duration of response and objective disease response (ORR, CR + CRu + PR). The complete response rate was 15% and overall response rate was 25%. Similar complete response rates were observed by the IRC across the 3 major PTCL subtypes (NOS, AITL, and ALK-1 negative ALCL). Median time to objective response was 1.8 months (~2 cycles) for the 33 patients who achieved CR, CRu or PR and was 3.7 months (~4 cycles) for the 19 patients with complete response. The responses in 11 of the 19 patients achieving CR and CRu were known to exceed 9.2 months; the follow-up on the remaining 8 patients was discontinued prior to 9.2 months.

Safety

The safety of Istodax® (romidepsin) was evaluated in 185 patients with CTCL in 2 single arm clinical studies in which patients received a starting dose of 14 mg/m2. The mean duration of treatment in these studies was 5.6 months (range: <1 to 83.4 months). Infections were the most common type of serious adverse event reported in both studies with 8 patients (8%) in Study 1 and 26 patients (31%) in Study 2 experiencing a serious infection. Serious adverse reactions reported in > 2% of patients in Study 1 were sepsis and pyrexia (3%). In Study 2, serious adverse reactions in > 2% of patients were fatigue (7%), supraventricular arrhythmia, central line infection, neutropenia (6%), hypotension, hyperuricemia, edema (5%), ventricular arrhythmia, thrombocytopenia, nausea, leukopenia, dehydration, pyrexia, aspartate aminotransferase increased, sepsis, catheter related infection, hypophosphatemia and dyspnea (4%). Most deaths were due to disease progression. In Study 1, there were two deaths due to cardiopulmonary failure and acute renal failure. In Study 2, there were six deaths due to infection (4), myocardial ischemia, and acute respiratory distress syndrome. Discontinuations Discontinuation due to an
adverse event occurred in 21% of patients in Study 1 and 11% in Study 2. Discontinuations occurring in at least 2% of patients in either study included infection, fatigue, dyspnea, QT prolongation, and hypomagnesemia.

**Zolinza® (vorinostat)**

The efficacy and safety of Zolinza® (vorinostat) was established in two nonrandomized, uncontrolled, open-label studies in adult patients with treatment refractory CTCL. A response rate (defined as at least 50% improvement in cutaneous manifestations) of approximately 30% was observed in patients receiving the approved dosing regimen of 400 mg once daily. Median time to response was <3 months, median response duration was approximately 3-4 months, and median time to progression was approximately 6-7 months.

**Beleodaq® (belinostat)**

**Efficacy**

Beleodaq® (belinostat) is a histone deacetylase (DHAC) inhibitor indicated for the treatment of patients with relapsed or refractory peripheral T-cell lymphoma (PTCL). It received accelerated approval based on the results of 1 single-arm, phase 2 study using tumor response rate as the primary outcome measure. An improvement in survival or disease-related symptoms has not been established. The pivotal trial has not been published.

In all evaluable patients (n=120) treated with belinostat, the overall response rate per independent central review using the International Workshop Group (IWG) criteria was 25.8% (n=31), with rates of 23% for PTCL not otherwise specified (NOS) and 46% for angioimmunoblastic T-cell lymphoma (AITL), the 2 largest subtypes enrolled. Further, the response rate was higher (28%) in patients with baseline platelet counts at or above 100,000/µL and lower (15%) in those with baseline platelet counts below 100,000/µL. The median duration of response based on the first date of response to disease progression or death was 8.4 months (95% confidence interval [CI], 4.5 to 29.4). Of the responders, the median time to response was 5.6 weeks (range, 4.3-50.4 weeks).
Safety

Sixty-one patients (47.3%) in the pivotal study experienced serious adverse reactions (grade 3 or higher) while taking belinostat or within 30 days after their last dose. The most common serious adverse reactions (>2%) were pneumonia, pyrexia, infection, anemia, increased creatinine, thrombocytopenia, and multiorgan failure. One treatment-related death associated with hepatic failure was reported in the study. Twenty-five patients (19.4%) in the pivotal study discontinued treatment with belinostat due to adverse reactions. Dose adjustments were made in 12%.

Valchlor® (mechlorethamine)

Efficacy

Valchlor® (mechlorethamine) is an alkylating agent indicated for the topical treatment of Stage IA and IB mycosis fungoides-type cutaneous T-cell lymphoma in patients who have received prior skin-directed therapy. The efficacy of Valchlor® (mechlorethamine) was assessed in a randomized, multicenter, observer-blind, active-controlled, non-inferiority clinical trial of 260 patients with Stage IA, IB, and IIA mycosis fungoides-type cutaneous T-cell lymphoma (MF-CTCL) who had received at least one prior skin-directed therapy. Qualifying prior therapies included topical corticosteroids, phototherapy, Targretin® gel, and topical nitrogen mustard. Patients were not required to be refractory to or intolerant of prior therapies. Study drug was to be applied topically on a daily basis for 12 months. Concomitant use of topical corticosteroids was not permitted during the study. Dosing could be suspended or continued with reduced frequency for dermatitis. The mean daily usage of Valchlor® gel was 2.8 g (1 to 2 tubes per month). The maximum daily usage was 10.5 g (5 to 6 tubes per month). Patients were evaluated for a response on a monthly basis for the first 6 months and then every 2 months for the last 6 months using the Composite Assessment of Index Lesion Severity (CAILS) score. The CAILS score is obtained by adding the severity score of each of the following categories for up to 5 index lesions: erythema, scaling, plaque elevation, and surface area. Severity was graded from 0 (none) to 8 (severe) for erythema and scaling; 0 to 3 for plaque elevation; and 0 to 9 for surface area. A response was defined as greater than or equal to 50% reduction in baseline CAILS score which was confirmed at the next visit at least 4 weeks later. A complete response was defined as a confirmed CAILS score of 0. Non-inferiority was considered to have been demonstrated if the lower bound of the 95% confidence interval (CI) around the ratio of response rates (Valchlor®/Comparator) was greater than or equal to 0.75. Patients were also evaluated using the Severity Weighted Assessment Tool (SWAT). The SWAT score is derived by measuring each involved area as a percentage of total body surface area (%BSA) and multiplying it by a severity weighting factor (1=patch, 2=plaque, 3=tumor or ulcer). A response was defined as greater than...
or equal to 50% reduction in baseline SWAT score which was confirmed at the next visit at least 4 weeks later. Sixty percent (60%) of the patients on the Valchlor® arm and 48% of patients on the comparator arm achieved a response based on the CAILS score. Valchlor® was noninferior to the comparator based on a CAILS overall response rate ratio of 1.24 (95% CI: 0.98, 1.58). Complete responses constituted a minority of the CAILS or SWAT overall responses. The onset of CAILS overall response for both treatment arms showed a wide range from 1 to 11 months.

## Safety

In the clinical trial, moderately-severe to severe skin-related adverse events were managed with treatment reduction, suspension, or discontinuation. Discontinuations due to adverse reactions occurred in 22% of patients treated with Valchlor® and 18% of patients treated with the comparator. Sixty-seven percent (67%) of the discontinuations for adverse reactions occurred within the first 90 days of treatment. Temporary treatment suspension occurred in 34% of patients treated with Valchlor® and 20% of patients treated with the comparator. Reductions in dosing frequency occurred in 23% of patients treated with Valchlor® and 12% of patients treated with the comparator. Reductions in hemoglobin, neutrophil count, or platelet count occurred in 13% of patients treated with Valchlor® and 17% treated with Comparator. Systemic exposure was undetectable after topical administration of Valchlor® to patients. Blood samples were analyzed from 16 and 15 patients following treatment with Valchlor® (mechlorethamine gel 0.016%) and an identical formulation consisting of mechlorethamine 0.032% w/w, respectively. For patients who received mechlorethamine 0.016%, samples were collected to measure mechlorethamine concentrations prior to dosing, on day 1, and at the first month visit. Following the topical administration of mechlorethamine 0.016%, there were no detectable plasma mechlorethamine concentrations observed in any of the patients. Patients who received mechlorethamine 0.032% had no measurable concentrations of mechlorethamine or half-mustard after 2, 4, or 6 months of treatment.

For more details, please see package insert for mechlorethamine.

## 2013 Update

A literature search for new publications from 01/01/2012 to 04/30/2013 did not reveal new evidence that would require changes to this policy. The policy was compared with current NCCN guideline recommendations and found to be consistent. An updated systematic review of treatment for mycosis fungoides/Sezary syndrome was published by the Cochrane Skin Group in September 2012.
2014 Update

A literature search for new publications from 01/01/2013 to 10/31/2014 did not reveal new evidence that would require changes to the drugs previously in this policy. Added Beleodaq® (belinostat), recently approved by FDA with medically necessary indications consistent with current NCCN guideline recommendations.

2015 Update

Valchlor® (mechlorethamine) was added to the policy on 07/27/2015, per the NCCN guideline recommendations.

2016 Update

Safety and efficacy reorganized to be under each relevant drug. No major NCCN guideline changes at this time.

2018 Update

A primary literature search from 04/11/2017 to 03/13/2018 did not reveal new evidence that would require change in this policy. No major NCCN guideline changes at this time.

References


25. Istodax® (romidepsin) prescribing information. Celgene Corporation; Summit, NJ. September 2011
Data on file, Celgene Corporation.


### History

<table>
<thead>
<tr>
<th>Date</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>05/10/11</td>
<td>Add to Prescription Drug Section - New Policy.</td>
</tr>
<tr>
<td>04/10/12</td>
<td>Replace policy. Policy updated with romidepsin (Istodax®) and a new medically necessary policy statement for the treatment of mycosis fungoides/Sezary syndrome for patients refractory to at least one prior systemic therapy. Description, Policy Guidelines and Rationale also updated.</td>
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<tr>
<td>05/28/13</td>
<td>Replace policy. Policy reviewed with literature search/ reference added. No change in policy statement.</td>
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<tr>
<td>12/08/14</td>
<td>Annual review. Medically necessary policy statement added for belinostat, recently approved by the FDA, per NCCN guideline recommendations; approved by P&amp;T December 2014. Literature review performed.</td>
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<tr>
<td>06/18/15</td>
<td>Update Related Policies. Change title to 8.01.36</td>
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<tr>
<td>08/11/15</td>
<td>Annual Review. Added a new agent Valchlor® to the existing policy. Updated Policy Guidelines and Rationale sections accordingly.</td>
</tr>
<tr>
<td>01/19/16</td>
<td>Coding update. New HCPCS code J9032, effective 1/1/16, added to policy. Minor edit to correct spelling and punctuation.</td>
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<tr>
<td>02/23/16</td>
<td>Coding update. Add J9315.</td>
</tr>
<tr>
<td>05/01/17</td>
<td>Annual Review, changes approved April 11, 2017. Criteria for topical bexarotene gel have been added. Also, statement outlining the length of therapy for initial and subsequent approval has been added to the policy.</td>
</tr>
<tr>
<td>10/24/17</td>
<td>Policy moved to new format; no change to policy statements.</td>
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</tbody>
</table>
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U.S. Department of Health and Human Services
200 Independence Avenue SW, Room 509F, HHH Building
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Deutsche (German):

Hmoob (Hmong):

Ilokano (Ilocano):
Daytoby a Pakdaak ket naglaon iti Napateg nga Impormasion. Daytoby a pakdaak mabalbin nga adda ket naglaon iti napateg nga impormasion maipanggpe iti aplikasyonono yewno coverage babaen iti Premera Blue Cross. Daytoby ket mabalbin dagiti importante a pelsa iti daytoby a pakdaak. Mabalbin nga adda rumbeng ng a aramidenyo nga addang sakkay dagiti partikular a naituding nga aildaw tapno mapagtalaindyo ti coverage ti salun-ayno yewno tulong kadagiti gastos. Adda karbenganyo a mangala iti daytoby nga impormasion ken tulong ti bukodyo a pagasasa nga awan ti bayadanyo. Tumawag ti numero nga 800-722-1471 (TTY: 800-842-5357).

Italiano (Italian):
Premera Blue Cross was not able to verify you in a timely manner.

If you have an unlisted phone number, please contact Premera Blue Cross at 800-722-1471 (TTY: 800-842-5357) for assistance.

If you do not have a phone number listed with Premera Blue Cross, you may be able to update your personal information online at premere.bcbs.com or call Premera Blue Cross at 800-722-1471 (TTY: 800-842-5357).

Please visit the website or call Premera Blue Cross for further information.

If you do not have internet access, you may contact Premera Blue Cross at 800-722-1471 (TTY: 800-842-5357) for assistance.

If you have questions about the notification, please call Premera Blue Cross at 800-722-1471 (TTY: 800-842-5357).

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