# SPECIALTY GUIDELINE MANAGEMENT

## TREANDA (bendamustine) BENDEKA (bendamustine)

### POLICY

#### I. INDICATIONS

The indications below including FDA-approved indications and compendial uses are considered a covered benefit provided that all the approval criteria are met and the member has no exclusions to the prescribed therapy.

- A. <u>FDA-Approved Indications<sup>1,2</sup></u>
  - 1. Chronic lymphocytic leukemia (CLL)
  - 2. Indolent B-cell non-Hodgkin lymphoma (NHL) that has progressed during or within six months of treatment with rituximab or a rituximab-containing regimen
- B. <u>Compendial Uses</u><sup>3,4</sup>
  - 1. Classical Hodgkin lymphoma (CHL)
  - 2. Multiple myeloma (MM)
  - 3. Non-Hodgkin lymphoma (NHL)
    - i. Adult T-cell leukemia/lymphoma (ATLL)
    - ii. Acquired immune deficiency syndrome (AIDS)-related B-cell lymphoma
    - iii. CLL/small lymphocytic lymphoma (SLL)
    - iv. Diffuse large B-cell lymphoma (DLBCL)
    - v. Follicular lymphoma
    - vi. Marginal zone lymphoma
      - a. Nodal marginal zone lymphoma
      - b. Gastric mucosa associated lymphoid tissue (MALT) lymphoma
      - c. Nongastric MALT lymphoma
      - d. Splenic marginal zone lymphoma
    - vii. Mantle cell lymphoma (MCL)
    - viii. Mycosis fungoides (MF)/Sezary syndrome (SS)
    - ix. Peripheral T-cell lymphoma (PTCL)
    - x. Primary cutaneous B-cell lymphoma
    - xi. Primary cutaneous CD30+ T-cell lymphoproliferative disorder
    - xii. Post-transplant lymphoproliferative disorders
  - 4. Waldenström's macroglobulinemia/lymphoplasmacytic lymphoma

All other indications are considered experimental/investigational and are not a covered benefit.

### II. CRITERIA FOR INITIAL APPROVAL

#### A. Non-Hodgkin lymphoma (NHL)<sup>1-4</sup>

Authorization of 12 months may be granted for treatment of NHL with any of the following subtypes:

1. Follicular lymphoma

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- 2. Chronic lymphocytic leukemia/small lymphocytic leukemia (CLL/SLL) without chromosome 17p deletion or TP53 mutation
- 3. Diffuse large B-cell lymphoma (DLBCL)
- 4. Adult T-cell leukemia/lymphoma (ATLL)
- 5. AIDS-related B-cell lymphoma
- 6. Marginal zone lymphoma
  - i. Nodal marginal zone lymphoma
  - ii. Gastric MALT lymphoma
  - iii. Nongastric MALT lymphoma
  - iv. Splenic marginal zone lymphoma
- 7. Mantle cell lymphoma (MCL)
- 8. Mycosis fungoides (MF)/Sezary syndrome (SS)
- 9. Peripheral T-cell lymphoma (PTCL)
- 10. Primary cutaneous B-cell lymphoma
- 11. Cutaneous anaplastic large cell lymphoma (ALCL)
- 12. Post-transplant lymphoproliferative disorders
- **B. Waldenström's macroglobulinemia/lymphoplasmacytic lymphoma<sup>3</sup>** Authorization of 12 months may be granted for treatment of Waldenström's macroglobulinemia/lymphoplasmacytic lymphoma.

## C. Multiple myeloma (MM)<sup>3</sup>

Authorization of 12 months may be granted for treatment of MM.

**D.** Classical Hodgkin lymphoma (CHL)<sup>3</sup> Authorization of 12 months may be granted for treatment of CHL.

## **III. CONTINUATION OF THERAPY**

All members (including new members) requesting authorization for continuation of therapy must meet all initial authorization criteria.

## IV. REFERENCES

- 1. Treanda [package insert]. North Wales, PA: Teva Pharmaceuticals USA, Inc.; December 2017.
- 2. Bendeka [package insert]. North Wales, PA: Teva Pharmaceuticals USA, Inc.; September 2017.
- 3. The NCCN Drugs & Biologics Compendium<sup>®</sup> © 2018 National Comprehensive Cancer Network, Inc. <u>https://www.nccn.org</u>. Accessed March 30, 2018.
- 4. Clinical Consult: CVS Caremark Clinical Programs Review. Focus on Hematology-Oncology Clinical Programs. June 2018

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