Medical Drug Clinical Criteria

Subject: Tevimbra (tislelizumab-jsgr)

 Document #:
 CC-0262
 Publish Date:
 04/01/2025

 Status:
 Revised
 Last Review Date:
 02/21/2025

Table of Contents

Overview Coding References

<u>Clinical Criteria</u> <u>Document History</u>

Overview

This document addresses the use of Tevimbra (tislelizumab-jsgr). Tevimbra is a programmed death receptor-1 (PD-1) blocking antibody. The FDA approved indication for Tevimbra includes use in the treatment of adult patients with unresectable or metastatic esophageal squamous cell carcinoma (ESCC) after prior systemic chemotherapy that did not include a PD-(L)1 inhibitor.

Definitions and Measures

ECOG or Eastern Cooperative Oncology Group Performance Status: A scale and criteria used by doctors and researchers to assess how an individual's disease is progressing, assess how the disease affects the daily living abilities of the individual, and determine appropriate treatment and prognosis. This scale may also be referred to as the WHO (World Health Organization) or Zubrod score which is based on the following scale:

- 0 = Fully active, able to carry on all pre-disease performance without restriction
- 1 = Restricted in physically strenuous activity but ambulatory and able to carry out work of a light or sedentary nature, for example, light house work, office work
- 2 = Ambulatory and capable of all self-care but unable to carry out any work activities. Up and about more than 50% of waking hours
- 3 = Capable of only limited self-care, confined to bed or chair more than 50% of waking hours
- 4 = Completely disabled. Cannot carry on any self-care. Totally confined to bed or chair
- 5 = Dead

Metastasis: The spread of cancer from one part of the body to another; a metastatic tumor contains cells that are like those in the original (primary) tumor and have spread.

Clinical Criteria

When a drug is being reviewed for coverage under a member's medical benefit plan or is otherwise subject to clinical review (including prior authorization), the following criteria will be used to determine whether the drug meets any applicable medical necessity requirements for the intended/prescribed purpose.

Tevimbra (tislelizumab-jsgr)

Requests for Tevimbra (tislelizumab-jsgr) may be approved if the following criteria are met:

- Individual has a diagnosis of metastatic esophageal squamous cell carcinoma (ESCC); (Label); AND
- II. Disease has progressed during or after first-line treatment for advanced unresectable/metastatic ESCC; AND
- III. Individual has a current Eastern Cooperative Oncology Group (ECOG) performance status of 0-1; AND
- IV. Individual is using as a single agent.

OR

- Individual has a diagnosis of gastric cancer (gastric or gastroesophageal junction adenocarcinoma) (Label);
 AND
- VI. Individual has either unresectable or metastatic HER2-negative disease; AND
- VII. Individual has a tumor which expresses PD-L1 (≥1); **AND**
- VIII. Individual is using as first-line therapy; AND
- IX. Individual has not received another anti-PD-1 or anti-PD-L1 agent or any other antibody or drug specifically targeting T-cell co-stimulation or checkpoint pathways; **AND**
- X. Individual is using in combination with platinum and fluoropyrimidine-based chemotherapy; AND
- XI. Individual has a current ECOG performance status of 0-1;

OR

- XII. Individual is using for Chronic lymphocytic leukemia/Small lymphocytic leukemia (CLL/SLL) with 17p deletion (NCCN 2A); AND
- XIII. Individual is using Tevimbra (tislelizumab-jsgr) in combination with Brukinsa (zanubrutinib).

Requests for Tevimbra (tislelizumab-jsgr) may not be approved for the following:

- I. When using for ESCC:
 - Individual has used two or more prior systemic treatments for advanced/metastatic unresectable ESCC; OR
 - B. Individual has uncontrollable pleural effusion, pericardial effusion, or ascites requiring frequent drainage; **OR**
 - Individual received prior therapies targeting programmed death 1 (PD-1) or programmed death ligand 1 (PD-L1); OR
 - D. Individual has active brain or leptomeningeal metastasis; OR
 - Individual has active autoimmune disease or history of autoimmune disease at high risk for relapse;
 OR
 - F. Individual has known history of, or any evidence of interstitial lung disease, non-infectious pneumonitis, pulmonary fibrosis diagnosed based on imaging or clinical findings, or uncontrolled systemic diseases, including diabetes, hypertension, acute lung disease, etc; **OR**
- II. When using for gastric cancer:
 - A. Individual has squamous cell or undifferentiated or other histological type gastric cancer; OR
 - B. Individual has active leptomeningeal disease or uncontrolled brain metastasis; OR
 - C. Individual has active autoimmune disease or history of autoimmune disease, or a medical condition requiring systemic corticosteroids or immunosuppressants; **OR**
- III. When the above criteria are not met, and for all other indications.

Coding

The following codes for treatments and procedures applicable to this document are included below for informational purposes. Inclusion or exclusion of a procedure, diagnosis or device code(s) does not constitute or imply member coverage or provider reimbursement policy. Please refer to the member's contract benefits in effect at the time of service to determine coverage or non-coverage of these services as it applies to an individual member.

HCPCS

J9329 Injection, tislelizumab-jsgr, 1mg [Tevimbra]

ICD-10 Diagnosis

C15.3-C15.9 Malignant neoplasm of esophagus
C16.0-C16.9 Malignant neoplasm of stomach
C83.00-C83.09 Small cell B-cell lymphoma
C83.30-C83.38 Diffuse large B-cell lymphoma

C83.398 Diffuse large B-cell lymphoma of other extranodal and solid organ sites

C91.10-C91.12 Chronic lymphocytic leukemia of B-cell type

Z85.01 Personal history of malignant neoplasm of esophagus

Document History

Revised: 02/21/2025 Document History:

- 02/21/2025 Select: Add new FDA indication for use in gastric cancer. Add NCCN recommendation for use in CLL/SLL with Brukinsa (zanubrutinib). Update may not be approved section. Coding Reviewed: Consolidated ICD-10-CM C15.3-C15.9 into one range and updated description. Added C16.0-C16.9, C83.00-C83.09, C83.30-C83.38, C83.398, C91.10-C91.12.
- 05/17/2024 Select Review: New criteria document for Tevimbra PA. Coding Reviewed: Added HCPCS J3590, J9999. Added All diagnoses pend. CMS Update: Remove HCPCS J3590 and J9999 and replace with J9329 and add ICD-10-CM C15.3, C15.4, C15.5, C15.8, C15.9, Z85.01.

References

- 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.; 2024; Updated periodically.
- 4. NCCN Clinical Practice Guidelines in Oncology™. © 2024 National Comprehensive Cancer Network, Inc. For additional information visit the NCCN website: http://www.nccn.org/index.asp. Accessed on December 27, 2024.
 - a. Chronic Lymphocytic Leukemia/Small Lymphocytic Lymphoma. V1.2025. Revised October 1, 2024.
 - b. Esophageal and esophagogastric junction cancers. V5.2024. Revised December 20, 2024.

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

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