# **CLINICAL POLICY Tebentafusp-tebn**



**Clinical Policy: Tebentafusp-tebn (Kimmtrak)** 

Reference Number: CP.PHAR.575

Effective Date: 05/2022 Last Review Date: 04/2023

Coding Implications
Revision Log

### **Description**

Tebentafusp-tebn (Kimmtrak®) is a bispecific gp100 peptide-HLA-directed CD3 T cell engager.

### **FDA** Approved Indication(s)

Kimmtrak is indicated for the treatment of HLA-A\*02:01-positive adult patients with unresectable or metastatic uveal melanoma.

# Policy/Criteria

Provider must submit documentation (such as office chart notes, lab results or other clinical information) supporting that member has met all approval criteria.

It is the policy of PA Health and Wellness <sup>®</sup> that Kimmtrak is **medically necessary** when the following criteria are met:

### I. Initial Approval Criteria

- A. Uveal Melanoma (must meet all):
  - 1. Diagnosis of unresectable or metastatic uveal melanoma;
  - 2. Prescribed by or in consultation with an oncologist;
  - 3. Age  $\geq$  18 years;
  - 4. Disease is HLA-A\*02:01-positive;
  - 5. Request meets one of the following (a or b):\*
    - a. Dose does not exceed 20 mcg (1 vial) on Day 1, 30 mcg (1 vial) on Day 8, 68 mcg (1 vial) on Day 15, and 68 mcg (1 vial) weekly thereafter;
    - b. Dose is supported by practice guidelines or peer-reviewed literature for the relevant off-label use (*prescriber must submit supporting evidence*).

**Approval duration: 6 months** 

# **B. Other diagnoses/indications:** Refer to PA.CP.PMN.53

### **II.** Continued Therapy

- **A.** Uveal Melanoma (must meet all):
  - 1. Currently receiving medication via PA Health & Wellness benefit and documentation supports positive response to therapy or the Continuity of Care policy (PA.LTSS.PHAR.01) applies;
  - 2. Member is responding positively to therapy;
  - 3. If request is for a dose increase, request meets one of the following (a or b):\*
    - a. New dose does not exceed 68 mcg (1 vial) weekly;
    - b. New dose is supported by practice guidelines or peer-reviewed literature for the relevant off-label use (*prescriber must submit supporting evidence*).

#### **Approval duration: 12 months**

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### **B.** Other diagnoses/indications (must meet 1 or 2):

- 1. Currently receiving medication via PA Health and Wellness benefit and documentation supports positive response to therapy or the Continuity of Care policy (PA.LTSS.PHAR.01) applies; or
- 2. Refer to PA.CP.PMN.53.

### III. Appendices/General Information

Appendix A: Abbreviation/Acronym Key

CRS: cytokine release syndrome FDA: Food and Drug Administration

*Appendix B: Therapeutic Alternatives* Not applicable.

Appendix C: Contraindications/Boxed Warnings

• Contraindication(s): none reported

• Boxed warning(s): cytokine release syndrome (CRS)

IV. Dosage and Administration

Indication	Dosing Regimen	Maximum Dose
Uveal melanoma	20 mcg on Day 1,	68 mcg /week
	30 mcg on Day 8,	
	68 mcg on Day 15, then	
	68 mcg once every week	
	thereafter	

### V. Product Availability

Injection: 100 mcg/0.5 mL vial

#### VI. References

- 1. Kimmtrak Prescribing Information. Conshohocken, PA: Immunocore Commercial Limited; November 2022. Available at: https://www.kimmtrak.com/. Accessed January 24, 2023.
- 2. Clinical Pharmacology [database online]. Tampa, FL: Elsevier, Inc.; 2023. Available at: http://www.clinicalkeys.com/pharmacology. Accessed January 24, 2023.
- 3. National Comprehensive Cancer Network. Melanoma: Uveal Version 2.2022 Available at: <a href="https://www.nccn.org/professionals/physician\_gls/pdf/uveal.pdf">https://www.nccn.org/professionals/physician\_gls/pdf/uveal.pdf</a>. Accessed January 242023.
- 4. National Comprehensive Cancer Network Drugs and Biologics Compendium. Available at www.nccn.org. Accessed January 24, 2023.

### **Coding Implications**

Codes referenced in this clinical policy are for informational purposes only. Inclusion or exclusion of any codes does not guarantee coverage. Providers should reference the most up-to-date sources of professional coding guidance prior to the submission of claims for reimbursement of covered services.

# CLINICAL POLICY Tebentafusp-tebn



HCPCS Codes	Description
J9274	Injection, tebentafusp-tebn, 1 mcg

Reviews, Revisions, and Approvals	Date	P&T Approval Date
Policy created.	04/2022	
Added HCPCS code [J9274].	01/2023	
2Q 2023 annual review: no significant changes; removed inactive	04/2023	
HCPCS codes C9095, J9999; references reviewed and updated.		