

| Vijoice (alpelisib) Effective 09/01/2025 | | | | | |
|---|--|---------------------------------------|----------------------|---------------------------------|--|
| Plan | ☐ MassHealth UPPL☒ Commercial/Exchange | | ⊠ Prior Authorizatio | | |
| Benefit | ☑ Pharmacy Benefit☐ Medical Benefit | | Program Type | ☐ Quantity Limit ☐ Step Therapy | |
| Specialty Limitations | This medication has been designated specialty and must be filled at a contracted specialty pharmacy. | | | | |
| | Medical and Specialty Medications | | | 3 | |
| Contact Information | All Plans | Phone: 877-519-1908 Fax: 855-540-3693 | | | |
| | Non-Specialty Medications | | | | |
| | All Plans | Pł | none: 800-711-4555 | Fax: 844-403-1029 | |

Overview

Vijoice is a kinase inhibitor indicated for the treatment of adult and pediatric patients 2 years of age and older with severe manifestations of PIK3CA-Related Overgrowth Spectrum (PROS) who require systemic therapy.

Coverage Guidelines

Exceptions

Authorization may be granted for members new to the plan within the past 90 days who are currently receiving treatment with requested medication, excluding when the product is obtained as samples or via manufacturer's patient assistance programs.

OR

Authorization may be granted when all of the following criteria are met:

- 1. Member has a diagnosis of PIK3CA-Related Overgrowth Spectrum (PROS)
- 2. Member is at least 2 years of age

N/A

- 3. The member has severe manifestations of disease and requires systemic therapy
- 4. Documented test confirming presence of PIK3CA mutation

Continuation of Therapy

Requests for reauthorization will be approved when all of the following criteria are met:

1. Attestation that the member has no evidence of unacceptable toxicity or disease progression while on the current regimen.

Limitations

- 1. Initial approvals will be granted for 6 months
- 2. Reauthorizations will be granted for 12 months.
- 3. The following quantity limitations apply:

| Drug Name and Dosage Form | Quantity Limit | |
|------------------------------|-------------------|--|
| Vijoice 50 mg, 125 mg tablet | 1 tablet per day | |
| Vijoice 250 mg tablet | 2 tablets per day | |
| Vijoice 50 mg granule | 1 packet per day | |

References

1. Vijoice [package insert]. East Hanover, NJ: Novartis Pharmaceuticals Corporation; April 2024.

Review History

09/21/2022 – Reviewed and created for July P&T. Effective 11/01/2022

08/14/2024 – Reviewed and updated at August P&T. Added Vijoice granule to the policy. Administrative update, specified quantity limitations. Effective 10/1/2024.

08/13/2025 – Reviewed at August P&T. No clinical changes. Effective 09/01/2025.

