Clinical Policy: Dordaviprone (Modeyso)

Reference Number: CP.PHAR.745
Effective Date: 12.01.25
Last Review Date: 11.26
Line of Business: Commercial, HIM/ICHRA, Medicaid Revision Log

See Important Reminder at the end of this policy for important regulatory and legal information.

Description

Dordaviprone (Modeyso™) is a protease activator.

FDA Approved Indication(s)

Modeyso is indicated for the treatment of adult and pediatric patients 1 year of age and older with diffuse midline glioma harboring an H3 K27M mutation with progressive disease following prior therapy.

This indication is approved under accelerated approval based on response rate and duration of response. Continued approval for this indication may be contingent upon verification and description of clinical benefit in a confirmatory trial(s).

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 health plans affiliated with Centene Corporation® that Modeyso is **medically necessary** when the following criteria are met:

I. Initial Approval Criteria

A. Diffuse Midline Glioma (must meet all)**:**

1. Diagnosis of diffuse midline or high-grade (*see Appendix D*) glioma;
2. Prescribed by or in consultation with an oncologist;
3. Age ≥ 1 year;
4. Disease has both of the following characteristics (a and b):
   a. Presence of H3 K27M mutation;
   b. Recurrent or progressive following prior therapy (includes radiation therapy; see *Appendix B* for examples of pharmacotherapies);
5. For Modeyso requests, member must use dordaviprone, if available, unless contraindicated or clinically significant adverse effects are experienced;
6. Request meets one of the following (a or b):*
   a. Dose does not exceed one of the following (i or ii):
      i. Adults: 625 mg per week;
      ii. Pediatrics (1, 2, 3, 4, or 5):
         1) Weight 10 kg to < 12.5 kg: 125 mg per week;
         2) Weight 12.5 kg to < 27.5 kg: 250 mg per week;
         3) Weight 27.5 kg to < 42.5 kg: 375 mg per week;

4) Weight 42.5 kg to < 52.5 kg: 500 mg per week;
5) Weight ≥ 52.5 kg: 625 mg per week;

b. Dose is supported by practice guidelines or peer-reviewed literature for the relevant off-label use (*prescriber must submit supporting evidence*).

**Prescribed regimen must be FDA-approved or recommended by NCCN*

Approval duration: 12 months

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

1. If this drug has recently (within the last 6 months) undergone a label change (e.g., newly approved indication, age expansion, new dosing regimen) that is not yet reflected in this policy, refer to one of the following policies (a or b):
   a. For drugs on the formulary (commercial, health insurance marketplace/ICHRA) or PDL (Medicaid), the no coverage criteria policy for the relevant line of business: CP.CPA.190 for commercial, HIM.PA.33 for health insurance marketplace/ICHRA, and CP.PMN.255 for Medicaid; or
   b. For drugs NOT on the formulary (commercial, health insurance marketplace/ICHRA) or PDL (Medicaid), the non-formulary policy for the relevant line of business: CP.CPA.190 for commercial, HIM.PA.103 for health insurance marketplace/ICHRA, and CP.PMN.16 for Medicaid; or
2. If the requested use (e.g., diagnosis, age, dosing regimen) is NOT specifically listed under section III (Diagnoses/Indications for which coverage is NOT authorized) AND criterion 1 above does not apply, refer to the off-label use policy for the relevant line of business: CP.CPA.09 for commercial, HIM.PA.154 for health insurance marketplace/ICHRA, and CP.PMN.53 for Medicaid.

II. Continued Therapy

A. Diffuse Midline Glioma (must meet all)**:**

1. Currently receiving medication via Centene benefit, or documentation supports that member is currently receiving Modeyso for a covered indication and has received this medication for at least 30 days;
2. Member is responding positively to therapy;
3. For Modeyso requests, member must use dordaviprone, if available, unless contraindicated or clinically significant adverse effects are experienced;
4. If request is for a dose increase, request meets one of the following (a or b):*
   a. New dose does not exceed one of the following (i or ii):
      i. Adults: 625 mg per week;
      ii. Pediatrics (1, 2, 3, 4, or 5):
         1) Weight 10 kg to < 12.5 kg: 125 mg per week;
         2) Weight 12.5 kg to < 27.5 kg: 250 mg per week;
         3) Weight 27.5 kg to < 42.5 kg: 375 mg per week;
         4) Weight 42.5 kg to < 52.5 kg: 500 mg per week;
         5) Weight ≥ 52.5 kg: 625 mg per week;
   b. New dose is supported by practice guidelines or peer-reviewed literature for the relevant off-label use (*prescriber must submit supporting evidence*).

**Prescribed regimen must be FDA-approved or recommended by NCCN*

Approval duration: 12 months

B. Other diagnoses/indications (must meet 1 or 2)**:**
1. If this drug has recently (within the last 6 months) undergone a label change (e.g., newly approved indication, age expansion, new dosing regimen) that is not yet reflected in this policy, refer to one of the following policies (a or b):
   a. For drugs on the formulary (commercial, health insurance marketplace/ICHRA) or PDL (Medicaid), the no coverage criteria policy for the relevant line of business: CP.CPA.190 for commercial, HIM.PA.33 for health insurance marketplace/ICHRA, and CP.PMN.255 for Medicaid; or
   b. For drugs NOT on the formulary (commercial, health insurance marketplace/ICHRA) or PDL (Medicaid), the non-formulary policy for the relevant line of business: CP.CPA.190 for commercial, HIM.PA.103 for health insurance marketplace/ICHRA, and CP.PMN.16 for Medicaid; or
2. If the requested use (e.g., diagnosis, age, dosing regimen) is NOT specifically listed under section III (Diagnoses/Indications for which coverage is NOT authorized) AND criterion 1 above does not apply, refer to the off-label use policy for the relevant line of business: CP.CPA.09 for commercial, HIM.PA.154 for health insurance marketplace/ICHRA, and CP.PMN.53 for Medicaid.

III. Diagnoses/Indications for which coverage is NOT authorized:

A. Non-FDA approved indications, which are not addressed in this policy, unless there is sufficient documentation of efficacy and safety according to the off label use policies – CP.CPA.09 for commercial, HIM.PA.154 for health insurance marketplace/ICHRA, and CP.PMN.53 for Medicaid or evidence of coverage documents.

IV. Appendices/General Information

Appendix A: Abbreviation/Acronym Key
FDA: Food and Drug Administration
NCCN: National Comprehensive Cancer Network

Appendix B: Therapeutic Alternatives
This table provides a listing of preferred alternative therapy recommended in the approval criteria. The drugs listed here may not be a formulary agent for all relevant lines of business and may require prior authorization.

Drug Name	Dosing Regimen	Dose Limit/ Maximum Dose
Examples of pharmacotherapies for adult gliomas (not mutation specific): temozolomide, PCV (procarbazine/lomustine/vincristine), lomustine, carmustine, bevacizumab, etoposide, carboplatin, cisplatin	Varies	Varies
Examples of pharmacotherapies for H3 K27-altered pediatric diffuse midline glioma: temozolomide, lomustine	Varies	Varies

Therapeutic alternatives are listed as Brand name® (generic) when the drug is available by brand name only and generic (Brand name®) when the drug is available by both brand and generic.

Appendix C: Contraindications/Boxed Warnings
None reported

Appendix D: General Information
- The National Comprehensive Cancer Network (NCCN) recommends the use of Modeyso in pediatric diffuse high-grade glioma and adult high-grade glioma. Per NCCN,
  - In pediatrics, diffuse high-grade glioma excludes: IDH-mutant and 1p/19q co-deleted oligodendroglioma and IDH-mutant astrocytoma.
  - In adults, high-grade glioma includes: H3-mutated high-grade glioma, high-grade astrocytoma with piloid features, and pleomorphic xanthoastrocytoma WHO grade 3.

V. Dosage and Administration

Indication	Dosing Regimen	Maximum Dose
Diffuse midline glioma	Adults: 625 mg PO once weekly Pediatrics: • Weight 10 kg to < 12.5 kg: 125 mg PO once weekly • Weight 12.5 kg to < 27.5 kg: 250 mg PO once weekly • Weight 27.5 kg to < 42.5 kg: 375 mg PO once weekly • Weight 42.5 kg to < 52.5 kg: 500 mg PO once weekly • Weight ≥ 52.5 kg: 625 mg PO once weekly	625 mg/week

VI. Product Availability
Capsule: 125 mg

VII. References
1. Modeyso Prescribing Information. Palo Alto, CA: Jazz Pharmaceuticals, Inc.; January 2026. Available at: www.modeyso.com. Accessed July 13, 2026.
2. National Comprehensive Cancer Network Drugs and Biologics Compendium. Available at: http://www.nccn.org/professionals/drug_compendium. Accessed July 27, 2026.
3. National Comprehensive Cancer Network. Central Nervous System Cancers Version 3.2026. Available at: https://www.nccn.org/professionals/physician_gls/pdf/cns.pdf. Accessed July 27, 2026.
4. National Comprehensive Cancer Network. Pediatric Central Nervous System Cancers Version 1.2026. https://www.nccn.org/professionals/physician_gls/pdf/ped_cns.pdf. Accessed July 27, 2026.

Reviews, Revisions, and Approvals	Date	P&T Approval Date
Policy created	08.13.25	11.25
4Q 2026 annual review: added option for use in high-grade glioma and added recurrent as an additional disease qualifier per NCCN; added ICHRA line of business; references reviewed and updated.	07.28.26	11.26

Important Reminder

This clinical policy has been developed by appropriately experienced and licensed health care professionals based on a review and consideration of currently available generally accepted standards of medical practice; peer-reviewed medical literature; government agency/program approval status; evidence-based guidelines and positions of leading national health professional organizations; views of physicians practicing in relevant clinical areas affected by this clinical policy; and other available clinical information. The Health Plan makes no representations and accepts no liability with respect to the content of any external information used or relied upon in developing this clinical policy. This clinical policy is consistent with standards of medical practice current at the time that this clinical policy was approved. "Health Plan" means a health plan that has adopted this clinical policy and that is operated or administered, in whole or in part, by Centene Management Company, LLC, or any of such health plan's affiliates, as applicable.

The purpose of this clinical policy is to provide a guide to medical necessity, which is a component of the guidelines used to assist in making coverage decisions and administering benefits. It does not constitute a contract or guarantee regarding payment or results. Coverage decisions and the administration of benefits are subject to all terms, conditions, exclusions, and limitations of the coverage documents (e.g., evidence of coverage, certificate of coverage, policy, contract of insurance, etc.), as well as to state and federal requirements and applicable Health Plan-level administrative policies and procedures.

This clinical policy is effective as of the date determined by the Health Plan. The date of posting may not be the effective date of this clinical policy. This clinical policy may be subject to applicable legal and regulatory requirements relating to provider notification. If there is a discrepancy between the effective date of this clinical policy and any applicable legal or regulatory requirement, the requirements of law and regulation shall govern. The Health Plan retains the right to change, amend or withdraw this clinical policy, and additional clinical policies may be developed and adopted as needed, at any time.

This clinical policy does not constitute medical advice, medical treatment, or medical care. It is not intended to dictate to providers how to practice medicine. Providers are expected to exercise professional medical judgment in providing the most appropriate care, and are solely responsible for the medical advice and treatment of members. This clinical policy is not intended to recommend treatment for members. Members should consult with their treating physician in connection with diagnosis and treatment decisions.

Providers referred to in this clinical policy are independent contractors who exercise independent judgment and over whom the Health Plan has no control or right of control. Providers are not agents or employees of the Health Plan.

This clinical policy is the property of the Health Plan. Unauthorized copying, use, and distribution of this clinical policy or any information contained herein are strictly prohibited. Providers, members, and their representatives are bound to the terms and conditions expressed herein through the terms of their contracts. Where no such contract exists, providers, members and their representatives agree to be bound by such terms and conditions by providing services to members and/or submitting claims for payment for such services.

Note:
For Medicaid members, when state Medicaid coverage provisions conflict with the coverage provisions in this clinical policy, state Medicaid coverage provisions take precedence. Please refer to the state Medicaid manual for any coverage provisions pertaining to this clinical policy.

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