

# Review at Launch for New to Market Medications (for Ohio Only)

**Policy Number:** CSOH2024D0060.A  
**Effective Date:** February 1, 2024

[➔ Instructions for Use](#)

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| Related Policy   |
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| <ul style="list-style-type: none"> <li><a href="#">Off-Label/Unproven/New FDA Indication Specialty Drug Treatment (for Ohio Only)</a></li> </ul> |

  

| Related Document   |
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| <ul style="list-style-type: none"> <li><a href="#">Review at Launch Medication List</a></li> </ul> |

## Application

This Medical Benefit Drug Policy only applies to the state of Ohio. Any requests for services that are stated as unproven or services for which there is a coverage or quantity limit will be evaluated for medical necessity using Ohio Administrative Code 5160-1-01.

## Coverage Rationale

**This drug policy applies to new medications that are:**

- U.S. Food and Drug Administration (FDA) approved; **and**
- Healthcare provider administered; **and**
- Reimbursable on a member’s medical benefit

**This policy does not apply to:**

- Medications used for the treatment of oncological conditions (these therapies are addressed by other policies/programs)
- Investigational/experimental medications
- Antibiotics/anti-infectives
- Nuclear pharmacy products (materials used in nuclear medicine procedures)
- Vaccines

All new medications that are identified as being subject to this policy will be placed on the [Review at Launch Medication List](#) and reviewed upon FDA approval.

Medications will be reviewed based on:

- Health plan benefits and whether the medication is a covered/reimbursable service; **and**
- Medical necessity

Medical necessity reviews will be conducted using the following:

- A UnitedHealthcare Pharmacy and Therapeutics (UHC P&T) approved medical benefit drug policy; **or**
- Medicaid state criteria as required; **or**
- **All** of the following:
  - FDA approved labeling, including but not limited to indication, patient age requirements, dosing recommendations, contraindications, and clinical trial inclusion criteria (ex. genetic testing, comorbid conditions); **and**

- Compendia (if available); **and**
- Current standard of care, as per evidenced based literature (if available)

The medications identified on the [Review at Launch Medication List](#) will be subject to this policy until such time that UnitedHealthcare determines pre-service reviews are no longer necessary or the drugs are added to the Prior Authorization List.

Claims submitted for a medication identified on the [Review at Launch Medication List](#) will be reviewed against health plan benefits and for medical necessity, as per the above.

## Applicable Codes

The following list(s) of procedure and/or diagnosis codes is provided for reference purposes only and may not be all inclusive. Listing of a code in this policy does not imply that the service described by the code is a covered or non-covered health service. Benefit coverage for health services is determined by federal, state, or contractual requirements and applicable laws that may require coverage for a specific service. The inclusion of a code does not imply any right to reimbursement or guarantee claim payment. Other Policies and Guidelines may apply.

| HCPCS Code | Description                       |
|------------|-----------------------------------|
| C9399      | Unclassified drugs or biologicals |
| J3490      | Unclassified drugs                |
| J3590      | Unclassified biologics            |

## Background

The Review at Launch program provides UnitedHealthcare the ability to review, evaluate, and implement programs for new to market medications. Additionally, it provides the opportunity to assess the coverage status of these new medications, and properly re-direct providers to State Medicaid Fee-For-Service programs when appropriate. The medications may be added to the Prior Authorization List once they have been evaluated by the UnitedHealthcare Pharmacy and Therapeutics Committee and a final utilization management strategy has been determined.

## References

1. AHFS Drug information [website]. Available at: <http://www.ahfsdruginformation.com/>. Accessed August 30, 2023.
2. Clinical Pharmacology [database online]. Tampa, FL: Gold Standard, Inc.; 2017. Available at: <http://www.goldstandard.com>. Accessed August 30, 2023.
3. Micromedex 2.0 [database online]. Truven Health Analytics, Inc. Greenwood Village, CO. Available at: <http://www.micromedexsolutions.com>. Accessed August 30, 2023.
4. UpToDate [database online]. Available at: <http://www.uptodate.com/>. Accessed August 30, 2023.
5. InterQual® [website]. Available at: <https://prod.cue4.com/help/InterQualOnline/BookViewHelp/content/>.

## Policy History/Revision Information

| Date       | Summary of Changes   |
|------------|--|
| 06/01/2024 | <b>Related Document</b> <ul style="list-style-type: none"> <li>● Updated <i>Review at Launch Medication List</i>; removed Winrevair™ (sotatercept-csrk)</li> </ul>         |
| 05/06/2024 | <b>Related Document</b> <ul style="list-style-type: none"> <li>● Updated <i>Review at Launch Medication List</i>; added Beqvez™ (fidanacogene elaparvovec-dzkt)</li> </ul> |
| 04/12/2024 | <b>Related Document</b> <ul style="list-style-type: none"> <li>● Updated <i>Review at Launch Medication List</i>; added Winrevair™ (sotatercept-csrk)</li> </ul>           |

| Date       | Summary of Changes   |
|------------|--|
| 04/01/2024 | <p><b>Related Policy</b></p> <ul style="list-style-type: none"> <li>Updated reference link to reflect current policy title for <i>Off-Label/Unproven/New FDA Indication Specialty Drug Treatment (for Ohio Only)</i></li> </ul> <p><b>Related Document</b></p> <ul style="list-style-type: none"> <li>Updated <i>Review at Launch Medication List</i>: <ul style="list-style-type: none"> <li>Added: <ul style="list-style-type: none"> <li>Tofidence™ (tocilizumab-bavi)</li> <li>Tyenne® (tocilizumab-aazg)</li> </ul> </li> <li>Removed (prior authorization requirements effective Apr. 1, 2024): <ul style="list-style-type: none"> <li>Adzynma (ADAMTS13, recombinant-krhn)</li> <li>Daxxify® (daxibotulinumtoxinA-lanm)</li> <li>Eylea® HD (aflibercept)</li> <li>Omvoh™ (mirikizumab-mrkz)</li> <li>Pombiliti™ (cipaglicosidase alfa)</li> </ul> </li> </ul> </li> </ul> |
| 02/01/2024 | <p><b>Template Update</b></p> <ul style="list-style-type: none"> <li>Created state-specific policy version</li> </ul> <p><b>Application</b></p> <ul style="list-style-type: none"> <li>Modified language to indicate this Medical Benefit Drug Policy only applies to the state of Ohio; any requests for services that are stated as unproven or services for which there is a coverage or quantity limit will be evaluated for medical necessity using <i>Ohio Administrative Code 5160-1-01</i></li> </ul> <p><b>Coverage Rationale</b></p> <ul style="list-style-type: none"> <li>Added language to indicate medical necessity reviews will be conducted using Medicaid state criteria as required</li> </ul> <p><b>Supporting Information</b></p> <ul style="list-style-type: none"> <li>Archived previous policy version CS2022D0060G</li> </ul>                           |

## Instructions for Use

This Medical Benefit Drug Policy provides assistance in interpreting UnitedHealthcare standard benefit plans. When deciding coverage, the federal, state (Ohio Administrative Code [OAC]), or contractual requirements for benefit plan coverage must be referenced as the terms of the federal, state (OAC), or contractual requirements for benefit plan coverage may differ from the standard benefit plan. In the event of a conflict, the federal, state (OAC), or contractual requirements for benefit plan coverage govern. Before using this policy, please check the federal, state (OAC), or contractual requirements for benefit plan coverage. UnitedHealthcare reserves the right to modify its Policies and Guidelines as necessary. This Medical Benefit Drug Policy is provided for informational purposes. It does not constitute medical advice.

UnitedHealthcare may also use tools developed by third parties, such as the InterQual® criteria, to assist us in administering health benefits. The UnitedHealthcare Medical Benefit Drug Policies are intended to be used in connection with the independent professional medical judgment of a qualified health care provider and do not constitute the practice of medicine or medical advice.