## MedStar Family Choice

CHOICE				
ADMINISTRATIVE POLICY AND PROCEDURE				
Policy #:	218			
Subject:	Pharmacy Authorization Process			
Section:	Pharmacy			
Initial Effective Date:	01/01/2018			
Revision Effective Date(s):	07/18, 07/19, 07/20, 07/21, 07/22, 07/23, 07/24			
Historical Revision Date(s):				
Review Effective Date(s):				
Historical Review Date(s):				
<b>Responsible Parties:</b>	Health Plan Pharmacist, P&T Committee			
Responsible Department(s):	Clinical Operations			
Regulatory References:	MDH Standards and Reporting Requirements of Drug Use Management Programs for MCOs March 2024 version, 2.12, 3.0. COMAR 10.67.09.04 I (d), COMAR 10.67.09.04(A)(3) NCQA 2023:UM 5C: 2, 4, 7, 8, UM 7G-I, UM 11B(4), UM 11E			
Approved:	AVP Clinical Operations	Chief Medical Officer		

- Purpose: To define a process to ensure that members receive medically necessary medication promptly when the medication(s) are non-formulary or formulary with a prior authorization requirement.
- Scope: MedStar Family Choice Maryland
- Policy: MedStar Family Choice follows standard processes for evaluating requests for medications requiring prior authorization in a timely fashion, consistent with MD COMAR and NCQA standards.

## **Definitions:**

**Request**: When a member, prescriber, or dispensing pharmacy staff asks for coverage of a specific pharmaceutical product.

• Includes concurrent and pre-service requests as defined by NCQA 2024 standards.

**Urgent Request**: A request for pharmaceutical services where application of the time frame for making routine or non-life-threatening care determinations:

- Could seriously jeopardize the life, health, or safety of the member or others, due to the member's psychological state, or
- In the opinion of a practitioner with knowledge of the member's medical or behavioral condition, would subject the member to adverse health consequences without the care or treatment that is the subject of the request.

**Retrospective or Post-Service Request**: A request for coverage of pharmaceutical services that has been received, such as if a member paid out-of-pocket for a prescription and is now seeking reimbursement.

Medical Reviewer: Medical Director or Health Plan Pharmacist

## Procedure:

- 1. A request for a medication authorization may be initiated by the prescriber, member, or the dispensing pharmacy staff via telephone, fax or on the MedStar Family Choice website.
  - 1.1. Requests may be submitted by telephone or fax using the corresponding submission form (see 1.2).
    - 1.1.1. Fax number: 410-933-2274, 410-350-7492, or
    - 1.1.2. Phone to 410-933-2200 or 800-905-1722
  - 1.2. Forms are located on the MedStar Family Choice pharmacy website: <u>https://www.medstarfamilychoice.com/maryland-providers/pharmacy-prescription-information</u>
    - 1.2.1. General Medication Prior Authorization form
    - 1.2.2. Hepatitis C Medication Prior Authorization form
    - 1.2.3. Opioid Prior Authorization form
  - 1.3. For phone requests, preauthorization staff will record the date and time of the request, the member's name, telephone number, prescriber's name, details about the medication being requested.
  - 1.4. Requests must include clinical documentation to support medical necessity.
  - 1.5. After-hour pharmacy requests:
    - 1.5.1. Requests submitted outside of the normal business hours of Monday through Friday, 8:30 am to 5 pm, will be reviewed by the on-call Medical Reviewer.
    - 1.5.2. A preauthorization staff member is also on-call to receive requests, enter the data into the clinical software system, and forward them to a Medical Reviewer to evaluate for a decision. The preauthorization staff will process Medical Reviewer decisions by entering any needed overrides into the PBM system and communicating the decision to the requestor, member, and/or pharmacy as needed.
- 2. Preauthorization staff will enter all requests into the clinical software system and forward them to a Medical Reviewer to evaluate and render a decision. If needed, the preauthorization staff may:
  - 2.1. Contact the prescriber to request incomplete or missing clinical information when not included with the initial request.

- 2.1.1. Preauthorization staff will make at least one attempt to request needed clinical information when omitted.
- 2.2. For formulary medication requiring Prior Authorization (PA) or Step Therapy (ST):
  - 2.2.1. Capture the approval criteria of the requested medication from the "Prior Authorization and Step Therapy Table" within the request.
  - 2.2.2. Collate any clinical information documented within the request that supports the PA or ST requirements.
  - 2.2.3. If PA or ST requirements are not met, redirection to a formulary alternative may be done only under the supervision of a Medical Reviewer.
- 2.3. Document if a requested medication is non-formulary.
  - 2.3.1. Redirection to a formulary alternative is done only by a Medical Reviewer.
- 2.4. Document if the prescriber is a network or non-network provider.
- 2.5. Preauthorization staff may obtain additional clinical information using available electronic health information resources to provide the Medical Reviewer information needed to evaluate medical necessity. Resources may include, but are not limited to:
  - 2.5.1. MedStar Family Choice's clinical software system,
  - 2.5.2. MedStar system EMR,
  - 2.5.3. PBM Claims reporting database.
- 2.6. Summarize associated clinical documentation then forward to Medical Reviewer.
- 3. The Medical Reviewer evaluates requests received in the clinical software system for medical necessity and clinical appropriateness.
  - 3.1. If additional clinical information is needed for evaluation:
    - 3.1.1. The Medical Reviewer may ask preauthorization staff to procure the needed information.
    - 3.1.2. The Medical Reviewer may contact the prescriber for additional information.
    - 3.1.3. The Medical Reviewer may access the Chesapeake Regional Information System (CRISP), *excluding* the of the Prescription Drug Monitoring Portal (PDMP) portion of the data base.
  - 3.2. The Medical Reviewer determines the medical necessity as described in pharmacy Policy 205: Non-Formulary Medications, Section 7.
  - 3.3. The Medical Reviewer makes a final decision to approve, deny, or otherwise action any request deemed to require clinical review.
    - 3.3.1. The timeline for rendering a decision is described in Section 7 of this policy.
    - 3.3.2. For cases where the full information is not available, the Medical Reviewer evaluates available information to make a final decision.
    - 3.3.3. Requests may be redirected by the Medical Reviewer or withdrawn if original requestor agrees.
    - 3.3.4. If the Medical Reviewer successfully redirects a request to a formulary alternative that requires PA or ST, the initial request may be converted to a request for the redirected medication.
      - 3.3.4.1. Preauthorization staff may update the request when the Medical Reviewer has successfully redirected the request.
- 4. The default length of the approval period is the duration requested by the provider. 4.1. Approval durations may have drug-specific limitations.

- 4.1.1. Maximum 6-month approval for controlled medications.
- 4.1.2. Maximum 12-month approval for non-controlled medications.
- 4.1.3. Length of the approval period may be shorter as identified on the PA and ST table.
- 4.2. Approval periods may differ for initial versus renewal authorizations.
- 4.3. Medical Reviewers may use their clinical judgement to render partial approvals or shorter approval duration when clinically appropriate.
- 4.4. MedStar Family Choice reserves the right to discontinue an existing authorization when the new authorization would create a duplication of therapy; this may result when the prescriber has discontinued the previously authorized medication.
- 5. Preauthorization staff process and complete notification for all completed requests.
  - 5.1. Notification of a decision complies with the timelines as described in Section 7 of this policy.
  - 5.2. Approved request processing:
    - 5.2.1. Preauthorization staff will enter an override in the Pharmacy Benefits Management (PBM) claims system so the prescription will adjudicate for the approved duration, and
    - 5.2.2. Notify the dispensing pharmacy to reprocess the claim, and
    - 5.2.3. Communicate the approval to the requestor by phone, fax, or electronic portal.
  - 5.3. Denied request processing:
    - 5.3.1. Preauthorization staff notify the member and prescriber of the denial in writing. The denial letter includes:
      - 5.3.1.1. The specific reason(s) for the denial, in easily understood language.
      - 5.3.1.2. A reference to the benefit provision, guideline, protocol, or other criterion upon which the denial decision is based.
      - 5.3.1.3. Formulary alternatives, if applicable.
      - 5.3.1.4. Directions to access the Formulary and/or PA & ST Table if applicable.
      - 5.3.1.5. Name and credentials of the Medical Reviewer.
      - 5.3.1.6. Option to discuss the denial with the Medical Reviewer, if desired.
      - 5.3.1.7. The process and timeline for initiating an appeal.
      - 5.3.1.8. Any additional information needed for the appeal.
      - 5.3.1.9. A statement that Members may obtain, upon request, a copy of the actual benefit provision, guideline, protocol, or other criterion on which the denial decision is based.
      - 5.3.1.10. Any forms required e.g., if requesting a Brand medication and a MedWatch form is required for approval, MedStar Family Choice will include the form with the denial.
    - 5.3.2. The appeal process is described in Member Services policies:
      - 5.3.2.1. Policy 301: Member Appeals; and
      - 5.3.2.2. Policy 307: Provider Appeals.
  - 5.4. Partially approved request notification follows processes described in Section 6.2 and Section 6.3 of this policy.

- 6. Pharmacy requests are processed in accordance with NCQA Standards, the Maryland Department of Health Standards and Reporting Requirements for Drug Use Management Programs for MCOs, and the Code of Maryland Regulations (COMAR).
  - 6.1. Timelines for decision-making and notification are described in Table 1 below.
  - 6.2. Requests confirmed to be clinically urgent will be prioritized for processing, and notification of the decision will occur no more than 24 hours from the date of receipt.
  - 6.3. Requesting prescribers may speak to a Medical Reviewer at any time during request processing.
  - 6.4. Members may initiate a retrospective coverage request up to 180 calendar days after the date of service.
    - 6.4.1. If request approval occurs within three (3) days of the date of service, the member will be instructed to return to the pharmacy for a refund.
    - 6.4.2. If request approval occurs more than three (3) days after the date of service, members will be instructed to mail their pharmacy and cash register receipts using the form and instructions described in Appendix I.
  - 6.5. Requests that are redirected to a formulary-preferred alternative, or otherwise determined to be withdrawn will be processed within 24 hours from the date of receipt.

Request Type	Timeline for UM Decision Making	Timeline for Notification	Notification Method	Who Must Be Notified
All Requests other than Retrospective or Post-service • Non-Urgent Preservice Requests • Urgent Requests* • Urgent concurrent requests	Within 24 hours of the receipt date of the Request for Authorization. MedStar Family Choice MD will approve, deny, or request further information. If further clinical information is not received: a decision is made within 24 hours of the date of receipt.	Notification of the decision within 24 hours of the receipt date. A decision is made within 24 hours of the receipt date regardless of whether clinical information is received	Notice by telephone or another telecommuni cation device. Electronic or written (required for denials)	Telephone or Other Telecommunication Device (required): - Requesting practitioner/provider Written (required for denials): - Requesting facility - Requesting physician or clinician - PCP -Member or Member's authorized representative
Retrospective/ Post-Service Pharmacy Requests	Within 30 calendar days of the receipt date of the request.	Electronic or written within 30 calendar days of the initial receipt date of the request.	Electronic or written	-Member or Member's representative (verbal approval or written denial) -Treating physician or clinician or requesting provider - PCP (denial only)

## Table 1. Pharmacy request timelines and notification processes.

\*Urgent requests will be processed as expediently as feasible, not to exceed 24 hours from date of receipt.

	<ul> <li>07/24:</li> <li>Moved P&amp;T Committee listing from "Responsible Department" to "Responsible Parties" section.</li> <li>Reformatted font and procedure to improve readability.</li> <li>Updated all NCQA, MCO Standards, and COMAR to current year references.</li> <li>Updated policy Approver titles and removed individual names.</li> </ul>
	<ul> <li>Incorporated Policy 212: Pharmacy Prior Authorization</li> <li>Reworded the Purpose statement for clarity and to remove references to policies that cite this policy as a reference.</li> <li>Clarified the process for redirecting to formulary preferred alternative medications, with emphasis that this may be done only by a Medical Reviewer or under the direct supervision of a Medical Reviewer.</li> </ul>
	<ul> <li>Added additional fax number for initiating an authorization request.</li> </ul>
	<ul> <li>Removed obsolete references to the 2017 Hepatitis-C PA timeline and all references, obsolete.</li> </ul>
	Added Appendix I: Prescription Reimbursement Claim Form     O7/23:
Summary of Changes:	<ul> <li>07/23:</li> <li>Responsible Parties changed to Health Plan Pharmacist</li> <li>Health Plan Pharmacist added with Medical Director throughout Policy</li> <li>Updated regulatory reference MDH Standards to March 2023</li> <li>Updated NCQA reference to 2023 Standards</li> <li>Updated Approved by to: Dr. Wills and C. Attia</li> <li>Updated COMAR references</li> <li>Removed Table: Medication Request from Patient, Prescriber or Pharmacy</li> </ul>
	<ul> <li>07/22</li> <li>Responsible Parties changed to Dr. Gregory Dohmeier</li> <li>Removed from Responsible Parties: Dr. Gerry and Dr. Toye</li> <li>Updated Regulatory Reference to April 2022 MDH Standards</li> <li>Updated NCQA Reference to 2022 Standards</li> <li>Updated tables to reflect changes to UM5C for pharmacy 24-hour response time</li> <li>Embedded MDH Memorandum 12/07/2017 expanded and concatenated to pdf</li> <li>07/21:</li> </ul>
	<ul><li>Updated NCQA Reference to reflect 2021 Standards.</li><li>Added Maryland to Scope.</li></ul>

I	
	<ul> <li>Changed Case Management to Clinical Operations in Responsible Departments.</li> </ul>
	07/20:
	<ul> <li>Updated Regulatory References to reflect COMAR</li> </ul>
	recodification and 2020 NCQA Standards.
	Added the following COMAR references found in Procedure
	Section to Regulatory References COMAR 10.67.11.04,
	COMAR 10.67.09.04(A)(3).
	Revised timeline for Urgent Concurrent requests from 72
	hours to 24 hours, regardless of the presence or absence of
	complete clinical information.
	07/19:
	<ul> <li>Updated NCQA Reference to reflect 2019 Standards.</li> </ul>
	<ul> <li>Removed "Maryland" from scope.</li> </ul>
	<ul> <li>Removed "A" from any policy reference, as applicable.</li> </ul>
	<ul> <li>Updated Urgent Concurrent timeframe to 72 hours from</li> </ul>
	previous 24 hours.
	07/18:
	<ul> <li>Procedure 1. Updated to reflect methods of making requests</li> </ul>
	(MFC PA form and MFC Non-Formulary Request form).
	• Procedure 2. Requests are sent to the medical director via the
	clinical software system.
	<ul> <li>Notification methods and who must be notified updated for all</li> </ul>
	urgent timeframes.
	<ul> <li>Urgent pre-service and non-urgent notifications were revised</li> </ul>
	to say that decision will occur within 24 hours of complete
	clinical.
	<ul> <li>Added "MDH Memorandum dated 12/07/2017: Re: Hepatitis C</li> </ul>
	Medication Approval Timeline" as a reference and embedded
	it at end of policy.
	<ul> <li>Modified Effective Date to Initial Effective Dates; added Historical Revision Dates and Revision Effective Dates; and</li> </ul>
	added Historical Review Dates and Review Effective Dates.
	01/18:
	New Policy.

Appendix I: Prescription Reimbursement Claim Form

