

## **Medical Care Subcommittee**

## Policy and Procedure for Prescription Drug Formulary Review

- 1. A) Ryan White Program Prescription Drug Formulary Review Request Form Completed and Sent to Recipient office for non-ART medications or
  - B) Request can be brought to the Recipient and then presented to the Medical Care Subcommittee.
- 2. New ADAP formulary approved ART medications will be automatically added to the Ryan White Program Prescription Drug Formulary.
- 3. Medical Care Subcommittee will be consulted when ADAP medications are either added or deleted for possible inclusion or exclusion from the Ryan White Part A Formulary.
- 4. Reviews request by the Medical Care Subcommittee will:
  - a. Conduct a literature review.
    - 1. Studies used must be of sufficient scientific rigor to ensure confidence in the claimed effects
    - 2. Study designs and measurements must reflect current scientific standards.
  - b. Evaluate and assess if drug/product is superior, inferior, equal to other therapies on the formulary, safety record of product, compliance evidence and economic considerations/impact including, but not limited to, moratorium restrictions that medications be either life saving or cost-effective.
  - c. Conduct the review, whenever possible, prior to the next Medical Care Subcommittee meeting.
- 5. Members of the Medical Care Subcommittee will complete a disclosure form (**Attachment 1**) once a year in January, new members upon joining and then in January. Conflicted members will recuse themselves from the vote.
- 6. Non-members who submit a formulary request must complete a disclosure form (Attachment 2) prior to the subcommittee voting.
- 7. Based on the literature review, evaluation and conclusions drawn the subcommittee will determine whether or not to recommend the medication/product. The subcommittee will also determine effective date of inclusion or removal, if a letter of medical necessity or a monitoring of the product is warranted. Upon completion of the vote the conflicted members may return.

MCSC Reviewed/Approved July 22, 2011/Revised January 25, 2013/Revised July 26, 2019/Revised October 23, 2020 and approved October 30, 2020

Date of Request: 11 8 2022	G-GC USE ONLY
Date of Request: 11/8/2022	2 Date Received
Request for: Addition Deletion	Date of First PUPAP Review Date of Approval HRSA Drug Code
(1) Generic/Proprietary name of drug product:  Methodore for opioid use disorder at	
(2) Specific formulation(s) considered:	
(3) Specific indications for use:  - priorid the disorder	
Please list other products currently in the formulary which are considered similar to the proposed addition/deletion:	
Should there be any restrictions on the use of this product?  Only for patients with a product?	
Please summarize your reasons and justification for this request. Provide appropriate references where applicable.  Comparentive Regulatoric Centers are the only licensed.  DTVs in Miami	
(7) I understand that this request will be considered at the next meeting of the Pharmaceutical Utilization Physicians Advisory Panel (PUPAP) or the Medi	
Print Name: Phone/Pager Clinic Site:	
Please forward this request to:	

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