## 4123-6-21.3 **Outpatient medication formulary.**

- (A) The administrator hereby adopts the formulary indicated in <u>the</u> appendix A-to this rule, developed with the recommendation of the bureau's pharmacy and therapeutics committee, effective <u>June 1, 2019 September 1, 2020</u>.
- (B) The Except as otherwise provided in paragraph (F) of this rule, the formulary indicated in the appendix A to this rule shall constitute the complete list of medications that are approved for reimbursement by the bureau for the treatment of an occupational a work related injury or disease in an allowed claim when dispensed to an injured worker by a registered pharmacist from an enrolled outpatient pharmacy provider. Except as otherwise provided in paragraph (F) of this rule, drugs not listed in the formulary are not eligible for reimbursement by the bureau.
- (C) The formulary indicated in <u>the appendix A</u>-to this rule also contains specific reimbursement, prescribing or dispensing restrictions that have been placed on the use of listed drugs. The formulary will be reviewed <del>annually</del> and updated as necessary. The most current version will be electronically published by the bureau.
- (D) Based upon current medical literature and generally accepted best clinical practices the bureau's pharmacy and therapeutics committee shall evaluate and make recommendations to the administrator regarding the addition, deletion, or modification of coverage of medications listed in the formulary. Requests for pharmacy and therapeutics committee action on a specific drug may be initiated by the bureau's administrator, chief of medical services, chief medical officer, or pharmacy director. The administrator will consider current medical literature and best practices and the recommendations of the bureau's pharmacy and therapeutics committee when making additions, deletions, or modifications of coverage of medications listed in the formulary.
- (E) The bureau shall develop policies to perform provide an expedited review process for clinically or therapeutically unique medications when necessary. The bureau shall also develop policies to address the timely review of new drug products.
- (F) Notwithstanding paragraph (B) of this rule, in cases of medical necessity supported by elinical medical documentation and evidence of need the bureau may, with prior authorization, reimburse for: new
  - (1) New drugs approved for use in the United States by the food and drug administration (FDA) on or after the effective date of the formulary, and for new indications approved by the FDA on or after the effective date of the formulary for existing drugs that are not on the formulary, with prior authorization, for a period not to exceed one hundred eighty days from the adjudication date of the first prescription for the requested drug.

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- (2) Antineoplastic drugs prescribed for treatment of an allowed cancer condition in a claim.
- (G) Notwithstanding <u>the appendix A-to this rule</u>, in cases of medical necessity supported by <u>clinical medical documentation</u> and evidence of need the bureau may, <u>with prior authorization</u>, reimburse for new dosage forms or strengths approved by the FDA on or after the effective date of the formulary for existing drugs that are on the formulary, <u>with prior authorization</u>, for a period not to exceed one hundred eighty days from the adjudication date of the first prescription for the requested drug.

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Effective: 9/1/2020

Five Year Review (FYR) Dates: 5/29/2020 and 05/01/2025

## CERTIFIED ELECTRONICALLY

Certification

08/07/2020

Date

Promulgated Under: 119.03

Statutory Authority: 4121.12, 4121.30, 4121.31, 4121.44,

4121.441, 4123.05, 4123.66

Rule Amplifies: 4121.12, 4121.121, 4121.44, 4121.441, 4123.66

Prior Effective Dates: 09/01/2011, 02/01/2012, 09/01/2012, 04/01/2013,

01/02/2014, 09/01/2014, 05/01/2015, 12/01/2015, 01/01/2017, 10/01/2017, 05/01/2018, 01/10/2019,

06/01/2019