

This document applies to the following:

Formulary	Applies
Standard Control (SF)	<input checked="" type="checkbox"/>
Standard Control – Choice (SCCF)	<input checked="" type="checkbox"/>
Preferred Drug Plan Design (PDPD)	<input type="checkbox"/>
Advanced Control Specialty (ACSF)	<input checked="" type="checkbox"/>
Advanced Control Specialty – Choice (ACSCF)	<input checked="" type="checkbox"/>
Managed Medicaid Template (MMT)	<input type="checkbox"/>
Marketplace (MF)	<input type="checkbox"/>
Aetna Small Group Affordable Care Act (SG ACA)	<input type="checkbox"/>
Aetna Health Exchange (AHE)	<input type="checkbox"/>
Aetna Individual Lives (IVL)	<input type="checkbox"/>
Value (VF)	<input checked="" type="checkbox"/>

Formulary	Applies
New to Market (NTM)	<input type="checkbox"/>
Standard Formulary Chart (SFC)	<input checked="" type="checkbox"/>
Basic Control Chart Preferred Drug Plan Design (BCC PDPD)	<input type="checkbox"/>
Advanced Control Specialty Formulary Chart (ACSFC)	<input checked="" type="checkbox"/>
Value Formulary Chart (VFC)	<input checked="" type="checkbox"/>
Medical Benefit	<input type="checkbox"/>
Medical Benefit: Advanced Biosimilars First	<input type="checkbox"/>
Medical Benefit: Managed Medicaid (MMMB)	<input type="checkbox"/>
Medicare Part B	<input type="checkbox"/>
Medicare Part B: Advanced Biosimilars First	<input type="checkbox"/>

Exceptions Criteria

Cuprimine, Syprine

This document informs prescribers of preferred products and provides an exception process for targeted products through prior authorization.

These criteria were developed to align with the following: Standard Control Formulary (SF), Standard Control Choice Formulary (SCCF), Advanced Control Specialty Formulary (ACSF), Advanced Control Specialty – Choice Formulary (ACSCF), Value Formulary (VF), Standard Formulary Chart (SFC), Advanced Control Specialty Formulary Chart (ACSFC) and Value Formulary Chart (VFC).

Plan Design Summary

This program applies to Cuprimine, Syprine products specified in this document. Coverage for targeted products is provided based on clinical circumstances that would exclude the use of the preferred product and may be based on previous use of a product. The coverage review process will ascertain situations where a clinical exception can be made. This program applies to all members requesting treatment with a targeted product.

Each referral is reviewed based on all utilization management (UM) programs implemented for the client.

Table. Cuprimine, Syprine Products

Medications considered formulary or preferred on your plan may still require a clinical prior authorization review.

	Product(s)
Preferred	<ul style="list-style-type: none"> • penicillamine (generic) • trientine (generic)
Targeted	<ul style="list-style-type: none"> • Cuprimine (penicillamine) • Syprine (trientine)

Exception Criteria

Cuprimine

Coverage for Cuprimine is provided when both of the following criteria are met:

- Member has failed treatment with the preferred generic product (penicillamine) due to an intolerable adverse event (e.g., rash, nausea, vomiting) and the adverse event was not an expected adverse event attributed to the active ingredient as described in the prescribing information (i.e., known adverse reaction for both the brand and generic medication)
- The adverse event is documented in member's chart. Submission of one of the following is required for approval:
 - Specific and detailed chart documentation including description, date and time, severity of the adverse event, dosage, duration of generic medication treatment, required intervention (if any), and relevant tests or laboratory data (if any).
 - MedWatch form of this trial and failure including the adverse reaction.

Syprine

Coverage for Syprine is provided when both of the following criteria are met:

- Member has failed treatment with the preferred generic product (trientine) due to an intolerable adverse event (e.g., rash, nausea, vomiting) and the adverse event was not an expected adverse event attributed to the active ingredient as described in the prescribing information (i.e., known adverse reaction for both the brand and generic medication)
- The adverse event is documented in member's chart. Submission of one of the following is required for approval:
 - Specific and detailed chart documentation including description, date and time, severity of the adverse event, dosage, duration of generic medication treatment, required intervention (if any), and relevant tests or laboratory data (if any).

Reference number(s)
4875-D

- MedWatch form of this trial and failure including the adverse reaction.

References

1. Cuprimine [package insert]. Bridgewater, NJ: Bausch Health US, LLC; October 2020.
2. Penicillamine [package insert]. Baudette, MN: Ani Pharmaceuticals Inc.; September 2021.
3. Syprine [package insert]. Bridgewater, NJ: Bausch Health US, LLC; September 2020.
4. Trientine [package insert]. Parsippany, NJ: Actavis Pharma Inc.; January 2022.