Clinical Policy: Cysteamine oral (Cystagon, Procysbi)
Reference Number: CP.PHAR.155
Effective Date: 02.16
Last Review Date: 05.20
Line of Business: HIM, Medicaid

See Important Reminder at the end of this policy for important regulatory and legal information.

Description
Cysteamine bitartrate (Cystagon®, Procysbi®) is a cysteine-depleting agent.

FDA Approved Indication
Cystagon and Procysbi are indicated for the treatment of nephropathic cystinosis. Cystagon is indicated for both children and adults, while Procysbi is indicated for patients 1 year of age and older.

Policy/Criteria
Provider must submit documentation (such as office chart notes, lab results or other clinical information) supporting that member has met all approval criteria.

It is the policy of health plans affiliated with Centene Corporation® that Cystagon and Procysbi are medically necessary when the following criteria are met:

I. Initial Approval Criteria
   A. Nephropathic Cystinosis (must meet all):
      1. Diagnosis of nephropathic cystinosis confirmed by one of the following (a, b, or c):
         a. Increased leukocyte cystine concentration (normal concentration: < 0.2 nmol half-cystine/mg protein);
         b. Cystinosin, lysosomal cystine transporter gene mutation;
         c. Corneal crystals on slit lamp examination;
      2. If Procysbi is requested, medical justification supports inability to use Cystagon (e.g., contraindication to excipients in Cystagon);
      3. Dose does not exceed 1.95 g per m² per day.
      Approval duration: 6 months

   B. Other diagnoses/indications
      1. Refer to the off-label use policy for the relevant line of business if diagnosis is NOT specifically listed under section III (Diagnoses/Indications for which coverage is NOT authorized): HIM.PHAR.21 for health insurance marketplace and CP.PMN.53 for Medicaid.

II. Continued Therapy
   A. Nephropathic Cystinosis (must meet all):
      1. Currently receiving medication via Centene benefit or member has previously met initial approval criteria;
2. Member is responding positively to therapy as evidenced by improvement in the leukocyte cystine concentration within the past 3 months;
3. If request is for a dose increase, new dose does not exceed 1.95 g per m² per day.

**Approval duration: 12 months**

**B. Other diagnoses/indications (must meet 1 or 2):**
1. Currently receiving medication via Centene benefit and documentation supports positive response to therapy.
   **Approval duration: Duration of request or 6 months (whichever is less); or**
2. Refer to the off-label use policy for the relevant line of business if diagnosis is NOT specifically listed under section III (Diagnoses/Indications for which coverage is NOT authorized): HIM.PHAR.21 for health insurance marketplace and CP.PMN.53 for Medicaid.

**III. Diagnoses/Indications for which coverage is NOT authorized:**
   **A.** Non-FDA approved indications, which are not addressed in this policy, unless there is sufficient documentation of efficacy and safety according to the off label use policy – HIM.PHAR.21 for health insurance marketplace and CP.PMN.53 for Medicaid, or evidence of coverage documents.

**IV. Appendices/General Information**

*Appendix A: Abbreviation/Acronym Key*
- FDA: Food and Drug Administration
- WBC: white blood cell

*Appendix B: Therapeutic Alternatives*
Not applicable

*Appendix C: Contraindications/Boxed Warnings*
- Contraindication(s): hypersensitivity to penicillamine or cysteamine.
- Boxed warning(s): none reported.

*Appendix D: General Information*
A clinical trial compared Cystagon and Procysbi in 43 (40 pediatric and 3 adult) patients with nephropathic cystinosis. Prior to randomization, patients were to be on a stable dose of Cystagon administered every six hours. This trial demonstrated that at steady-state, Procysbi administered every 12 hours was non-inferior to Cystagon administered every 6 hours with respect to the depletion of white blood cell (WBC) cystine concentrations. The least-square mean value of WBC cystine was 0.52 ± 0.06 nmol ½ cystine/mg protein after 12 hours under Procysbi and 0.44 ± 0.06 nmol ½ cystine/mg protein after 6 hours under Cystagon; a difference of 0.08 ± 0.03 nmol ½ cystine/mg protein (95.8% Confidence Interval = 0.01 to 0.15). The goal of cysteamine therapy is to lower WBC cystine levels.
V. Dosage and Administration

<table>
<thead>
<tr>
<th>Indication</th>
<th>Dosing Regimen</th>
<th>Maximum Dose</th>
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</table>
| Cystagon   | Initial: 1/4 to 1/6 of the maintenance dose  
Recommended maintenance dose:  
For age < 12 years: 1.30 g/m²/day given in four divided doses  
For age ≥ 12 years: 2.0 g/day in four divided doses | 1.95 g/m²/day |
| Procysbi   | Cysteamine-naïve patients:  
Initial: 1/4 to 1/6 of the maintenance dose  
Recommended maintenance dose: 1.3 g/m²/day given in two divided doses  
Switching from Cystagon: the starting total daily dose of Procysbi is equal to the previous total daily dose of Cystagon. Divide the total daily dose by two and administer every 12 hours. | 1.95 g/m²/day |

VI. Product Availability

<table>
<thead>
<tr>
<th>Drug</th>
<th>Availability</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cystagon</td>
<td>Capsule: 50 mg, 150 mg</td>
</tr>
</tbody>
</table>
| Procysbi | Delayed-release capsule: 25 mg, 75 mg  
Delayed-release oral granule packet: 75 mg, 300 mg |

VII. References


Reviews, Revisions, and Approvals

<table>
<thead>
<tr>
<th>Description</th>
<th>Date</th>
<th>P&amp;T Approval Date</th>
</tr>
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<tbody>
<tr>
<td>Policy split from CP.PHAR.48 LSD</td>
<td>01.16</td>
<td>02.16</td>
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<td>Policy converted to new template</td>
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<tr>
<td>Age restriction removed.</td>
<td>12.16</td>
<td>02.17</td>
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<tr>
<td>Additional diagnostic criteria added.</td>
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<tr>
<td>Reasons to discontinue added to continuation criteria.</td>
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<tr>
<td>Positive response to therapy added.</td>
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<tr>
<td>Background section converted to new template.</td>
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**Clinical Policy**

**Cysteamine oral**

<table>
<thead>
<tr>
<th>Reviews, Revisions, and Approvals</th>
<th>Date</th>
<th>P&amp;T Approval Date</th>
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<tbody>
<tr>
<td>Policy converted to newer template. Age restriction added. Reasons to discontinue removed from continuation criteria.</td>
<td>09.05.17</td>
<td>11.17</td>
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<tr>
<td>Q2 2018 annual review: no significant changes; HIM added; age restriction removed; added requirement of a prior trial of Cystagon for all Procysbi requests; added specific parameters for documenting a positive response to therapy, for reauthorization; references reviewed and updated.</td>
<td>02.25.18</td>
<td>05.18</td>
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<tr>
<td>2Q 2019 annual review: no significant changes; references reviewed and updated.</td>
<td>02.28.19</td>
<td>05.19</td>
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<tr>
<td>2Q 2020 annual review: no significant changes; references reviewed and updated.</td>
<td>02.05.20</td>
<td>05.20</td>
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**Important Reminder**

This clinical policy has been developed by appropriately experienced and licensed health care professionals based on a review and consideration of currently available generally accepted standards of medical practice; peer-reviewed medical literature; government agency/program approval status; evidence-based guidelines and positions of leading national health professional organizations; views of physicians practicing in relevant clinical areas affected by this clinical policy; and other available clinical information. The Health Plan makes no representations and accepts no liability with respect to the content of any external information used or relied upon in developing this clinical policy. This clinical policy is consistent with standards of medical practice current at the time that this clinical policy was approved. “Health Plan” means a health plan that has adopted this clinical policy and that is operated or administered, in whole or in part, by Centene Management Company, LLC, or any of such health plan’s affiliates, as applicable.

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**Note: For Medicaid members,** when state Medicaid coverage provisions conflict with the coverage provisions in this clinical policy, state Medicaid coverage provisions take precedence. Please refer to the state Medicaid manual for any coverage provisions pertaining to this clinical policy.

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