Clinical Policy: Gastrointestinal Agents: Opioid-Induced Constipation

Reference Number: OH.PHAR.PPA.57
Effective Date: 01/01/2020
Last Review Date: N/A
Line of Business: Medicaid

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

Description

GASTROINTESTINAL AGENTS: OPIOID-INDUCED CONSTIPATION AGENTS

<table>
<thead>
<tr>
<th>NO PA REQUIRED “PREFERRED”</th>
<th>STEP THERAPY REQUIRED “PREFERRED”</th>
<th>PA REQUIRED “NON-PREFERRED”</th>
</tr>
</thead>
<tbody>
<tr>
<td>BISACODYL (generic of Dulcolax®)</td>
<td>AMITIZA® capsule (lubiprostone)</td>
<td>RELISTOR® tablets and subcutaneous injection (methylaltrexone bromide)</td>
</tr>
<tr>
<td>CASANTHRANOL/DOCUSATE SODIUM (generic of Peri-Colace®)</td>
<td>MOVANTIK® tablets (naloxegol)</td>
<td>SYMPROIC® (naldemedine)</td>
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<tr>
<td>LACTULOSE (generic of Chronulac®)</td>
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<td></td>
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<tr>
<td>POLYETHYLENE GLYCOL (generic of Miralax®)</td>
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<tr>
<td>PSYLLIUM FIBER (e.g. Konsyl®)</td>
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<tr>
<td>SENNA (generic of Senokot®)</td>
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</tbody>
</table>

FDA Approved Indication(s): Varies by drug product, please see package insert; clinical pharmacology or other appropriate clinical reference.

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 Buckeye Health Plan, an affiliate of Centene Corporation® that Non-preferred Opioid-Induced Constipation agents are medically necessary when the following criteria are met:

I. Initial Approval Criteria

A. PA Required Non-Preferred Agents or Step Therapy agents

1. Prescribed indication is FDA-approved or supported by standard pharmacopeias
2. Age ≥ 18 years;
3. Approval requires a history of chronic pain requiring continuous opioid therapy for 12 weeks or longer. Electronic PA will approve with a history of 90 days of opioid therapy in the previous 90 days, in addition to trials of preferred products.
4. If medication requires step therapy (see above table) member must have had an inadequate clinical response to preferred alternatives, including a trial of no less than 14 day trial of at least two medications not requiring prior approval unless one of the following:

a. Allergy to medications not requiring prior approval
b. Contraindication to or drug interaction with medications not requiring prior approval
c. History of unacceptable/toxic side effects to medications not requiring prior approval

5. If medication is a non-preferred agent member must have had an inadequate clinical response to preferred alternatives, including a trial of no less than 14 day trial of at least two step therapy products medications unless one of the following:

a. Allergy to medications not requiring prior approval
b. Contraindication to or drug interaction with medications not requiring prior approval
c. History of unacceptable/toxic side effects to medications not requiring prior approval

Approval duration: 12 months.

II. Diagnoses/Indications for which coverage is NOT authorized:
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 – CP.PMN.53 for Medicaid or evidence of coverage documents.

III. Appendices/General Information

Appendix B: Therapeutic Alternatives
This table provides a listing of preferred alternative therapy recommended in the approval criteria. The drugs listed here may not be a formulary agent for all relevant lines of business and may require prior authorization.

**See above tables for preferred alternatives** Dosing varies by drug product. See FDA approved dosing and administration.

Therapeutic alternatives are listed as Brand name® (generic) when the drug is available by brand name only and generic (Brand name®) when the drug is available by both brand and generic.

Appendix C: Contraindications/Boxed Warnings
• See package insert; clinical pharmacology or other appropriate clinical reference
IV. Dosage and Administration: varies by drug product. See package insert; clinical pharmacology or other appropriate clinical reference for FDA approved dosing and administration.

V. Product Availability: See package insert; clinical pharmacology or other appropriate clinical reference for product availability.

VI. References. Refer to package insert.

<table>
<thead>
<tr>
<th>Reviews, Revisions, and Approvals</th>
<th>Date</th>
<th>P&amp;T Approval Date</th>
</tr>
</thead>
<tbody>
<tr>
<td>New policy created.</td>
<td>10.19</td>
<td>N/A</td>
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</tbody>
</table>

**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.

The purpose of this clinical policy is to provide a guide to medical necessity, which is a component of the guidelines used to assist in making coverage decisions and administering benefits. It does not constitute a contract or guarantee regarding payment or results. Coverage decisions and the administration of benefits are subject to all terms, conditions, exclusions and limitations of the coverage documents (e.g., evidence of coverage, certificate of coverage, policy, contract of insurance, etc.), as well as to state and federal requirements and applicable Health Plan-level administrative policies and procedures.

This clinical policy is effective as of the date determined by the Health Plan. The date of posting may not be the effective date of this clinical policy. This clinical policy may be subject to applicable legal and regulatory requirements relating to provider notification. If there is a discrepancy between the effective date of this clinical policy and any applicable legal or regulatory requirement, the requirements of law and regulation shall govern. The Health Plan retains the right to change, amend or withdraw this clinical policy, and additional clinical policies may be developed and adopted as needed, at any time.

This clinical policy does not constitute medical advice, medical treatment or medical care. It is not intended to dictate to providers how to practice medicine. Providers are expected to exercise
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