Clinical Policy: Cardiovascular Agents: Antiarrhythmics
Reference Number: OH.PHAR.PPA.30
Effective Date: 01-01-2020
Last Review Date: 10-2019
Line of Business: Medicaid

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

Description
The following are Antiarrhythmic agents requiring prior authorization:

<table>
<thead>
<tr>
<th>CARDIOVASCULAR AGENTS: ANTIARRHYTHMICS</th>
<th>NO PA REQUIRED “PREFERRED”</th>
<th>PA REQUIRED “NON-PREFERRED”</th>
</tr>
</thead>
<tbody>
<tr>
<td>AMIODARONE (generic of Cordarone®) 200mg</td>
<td></td>
<td>AMIODARONE 100mg, 400mg</td>
</tr>
<tr>
<td>DISOPYRAMIDE PHOSPHATE IR (generic of Norpace®)</td>
<td></td>
<td>MULTAQ® (dronedarone)</td>
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<tr>
<td>DISOPYRAMIDE PHOSPHATE ER (generic of Norpace® CR)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>FLECAINIDE (generic of Tambocor®)</td>
<td></td>
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<tr>
<td>MEXILITINE</td>
<td></td>
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<tr>
<td>PROPafenone (generic of Rythmol®)</td>
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<tr>
<td>PROPafenone ER (generic of Rythmol SR®)</td>
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<td></td>
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<tr>
<td>QUINIDINE GLUCONATE ER</td>
<td></td>
<td></td>
</tr>
<tr>
<td>QUINIDINE SULFATE</td>
<td></td>
<td></td>
</tr>
<tr>
<td>QUINIDINE SULFATE ER</td>
<td></td>
<td></td>
</tr>
<tr>
<td>TIKOSYN® (dofetilide)</td>
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<td></td>
</tr>
</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® that Amiodarone 100mg, 400mg, Mulltaq (dronedarone) is medically necessary when the following criteria are met:

I. Initial Approval Criteria
   A. Prescribed indication is FDA-approved or supported by standard pharmacopeias
      1. Member has failed therapeutic trials of 30 days with ONE (1) preferred medications unless one of the following (a,b, or c)
         a. Allergy to all medications not requiring prior approval
         b. Contraindication to all 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 policies – CP.PMN.53 for Medicaid or evidence of coverage documents.

III. Appendices/General Information
   Appendix A: Abbreviation/Acronym Key
   FDA: Food and Drug Administration
   PA: Prior Authorization
   ER: Extended Release

   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 table 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></td>
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</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
Clinical Policy: Cardiovascular Agents: Antiarrhythmics

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 professional medical judgment in providing the most appropriate care, and are solely responsible for the medical advice and treatment of members. This clinical policy is not intended to recommend treatment for members. Members should consult with their treating physician in connection with diagnosis and treatment decisions.

Providers referred to in this clinical policy are independent contractors who exercise independent judgment and over whom the Health Plan has no control or right of control. Providers are not agents or employees of the Health Plan.

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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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