Clinical Policy: Tasimelteon (Hetlioz)
Reference Number: CP.PMN.104
Effective Date: 02.01.17
Last Review Date: 08.19
Line of Business: Commercial, HIM, Medicaid

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

Description
Tasimelteon (Hetlioz®) is a melatonin receptor agonist.

FDA Approved Indication(s)
Hetlioz is indicated for treatment of non-24-hour sleep-wake disorder (non-24).

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 Hetlioz is medically necessary when the following criteria are met:

I. Initial Approval Criteria
   A. Non-24-Hour Sleep-Wake Disorder (must meet all):
      1. Diagnosis of non-24-hour sleep-wake disorder;
      2. Prescribed by or in consultation with a specialist in sleep disorders;
      3. Failure of melatonin and ramelteon (Rozerem®), unless contraindicated or clinically significant adverse effects are experienced;
         *Prior authorization may be required for ramelteon
      4. Member is completely blind (no light perception);
      5. Dose does not exceed 20 mg (1 capsule) per day.
   Approval duration:
   Medicaid/HIM – 6 months
   Commercial – Length of Benefit

   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): CP.CPA.09 for commercial, HIM.PHAR.21 for health insurance marketplace, and CP.PMN.53 for Medicaid.

II. Continued Therapy
   A. Non-24-Hour Sleep-Wake Disorder (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;
3. If request is for a dose increase, new dose does not exceed 20 mg (1 capsule) per day.
   **Approval duration:**
   Medicaid/HIM – 12 months
   Commercial – Length of Benefit

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): CP.CPA.09 for commercial, 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 policies – CP.CPA.09 for commercial, 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

   *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.

<table>
<thead>
<tr>
<th>Drug Name</th>
<th>Dosing Regimen</th>
<th>Dose Limit/Maximum Dose</th>
</tr>
</thead>
<tbody>
<tr>
<td>melatonin</td>
<td>5 to 10 mg PO QHS</td>
<td>N/A</td>
</tr>
<tr>
<td>Rozerem (ramelteon)</td>
<td>8 mg PO QHS</td>
<td>8 mg/day</td>
</tr>
</tbody>
</table>

   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*
   None reported

V. **Dosage and Administration**

<table>
<thead>
<tr>
<th>Indication</th>
<th>Dosing Regimen</th>
<th>Maximum Dose</th>
</tr>
</thead>
<tbody>
<tr>
<td>Non-24-hr-sleep-wake disorder</td>
<td>20 mg PO once daily at the same time each night before bedtime</td>
<td>20 mg/day</td>
</tr>
</tbody>
</table>

VI. **Product Availability**
Capsule: 20 mg
VII. References

Reviews, Revisions, and Approvals

<table>
<thead>
<tr>
<th>Reviews, Revisions, and Approvals</th>
<th>Date</th>
<th>P&amp;T Approval Date</th>
</tr>
</thead>
<tbody>
<tr>
<td>HIM: Policy created.</td>
<td></td>
<td>12.16</td>
</tr>
<tr>
<td>1Q18 annual review: Policies combined for HIM and commercial; Medicaid line of business was added to criteria; Added specialist requirement; Added trial and failure of melatonin; Removed diagnosis with “confirmed by at least 14 days of documentation of progressively shifting sleep-wake times” due to added specialist requirement; References reviewed and updated</td>
<td>11.20.17</td>
<td>02.18</td>
</tr>
<tr>
<td>1Q 2019 annual review: no significant changes; references reviewed and updated.</td>
<td>11.20.18</td>
<td>02.19</td>
</tr>
<tr>
<td>Added trial and failure of Rozerem including therapeutic alternatives table information; references reviewed and updated.</td>
<td>05.15.19</td>
<td>08.19</td>
</tr>
</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
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.

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Providers, members and their representatives are bound to the terms and conditions expressed
herein through the terms of their contracts. Where no such contract exists, providers, members
and their representatives agree to be bound by such terms and conditions by providing services to
members and/or submitting claims for payment for such services.

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.

For Health Insurance Marketplace members, when applicable, this policy applies only when
the prescribed agent is on your health plan approved formulary. Request for non-formulary drugs
must be reviewed using the formulary exception policy.

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