Clinical Policy: Bedaquiline (Sirturo)
Reference Number: CP.PMN.212
Effective Date: 09.04.18
Last Review Date: 02.20
Line of Business: HIM, Medicaid

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

Description
Bedaquiline (Sirturo®) is a diarylquinoline antimycobacterial drug.

FDA Approved Indication(s)
Sirturo is indicated as part of combination therapy in the treatment of adult and pediatric patients (5 years and older and weighing at least 15 kg) with pulmonary multi-drug resistant tuberculosis (MDR-TB). Reserve Sirturo for use when an effective treatment regimen cannot otherwise be provided.

Limitation(s) of use:
- Do not use Sirturo for the treatment of:
  - Latent infection due to *Mycobacterium tuberculosis*
  - Drug-sensitive tuberculosis
  - Extra-pulmonary tuberculosis
  - Infections caused by non-tuberculous mycobacteria
- The safety and efficacy of Sirturo in the treatment of HIV infected patients with MDR-TB have not been established as clinical data are limited.

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 Sirturo is medically necessary when the following criteria are met:

I. Initial Approval Criteria
   A. Multi-Drug Resistant Tuberculosis without Pretomanid (must meet all):
      1. Diagnosis of MDR-TB;
      2. Prescribed by or in consultation with an infectious disease specialist or a pulmonologist;
      3. Age ≥ 5 years;
      4. Prescribed in combination with at least 3 other anti-tuberculosis agents (*Appendix B*);
      5. Documented resistance to fluoroquinolones, unless contraindicated or clinically significant adverse effects are experienced;
      6. Dose does not exceed one of the following (a or b):
         a. Weight ≥ 30 kg: 400 mg per day for the first 2 weeks, followed by 200 mg three times per week;
b. Weight 15 to 29 kg: 200 mg per day for the first 2 weeks, followed by 100 mg three times per week.

Approval duration: 24 weeks

B. Multi-Drug Resistant Tuberculosis with Pretomanid (must meet all):
   1. Diagnosis of pulmonary MDR-TB or XDR-TB;
   2. Prescribed by or in consultation with an infectious disease specialist;
   3. Age ≥ 17 years;
   4. Prescribed in combination with pretomanid and linezolid;
      *Prior authorization may be required for pretomanid and linezolid.
   5. Documented resistance to fluoroquinolones, unless contraindicated or clinically significant adverse effects are experienced;
   6. Dose does not exceed 400 mg per day for the first 2 weeks, followed by 200 mg three times per week.

Approval duration: 6 months

C. 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. Multi-Drug Resistant Tuberculosis without Pretomanid (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 one of the following (a or b):
      a. Weight ≥ 30 kg: 200 mg three times per week;
      b. Weight 15 to 29 kg: 100 mg three times per week.

Approval duration: up to a total duration of 24 weeks

B. Multi-Drug Resistant Tuberculosis with Pretomanid (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. Member continues to receive pretomanid;
   4. Member has previously received at least 4 weeks of linezolid 1,200 mg per day;
   5. If request is for a dose increase, new dose does not exceed 200 mg three times per week.

Approval duration: up to a total treatment duration of 6 months (9 months if evidence of delayed culture conversion)
C. 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 policies – 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
   MDR-TB: multi-drug resistant tuberculosis
   XDR-TB: extensively drug resistant tuberculosis

   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>amikacin/kanamycin</td>
<td>15 mg/kg IM or IV QD or 25 mg/kg PO 3 times weekly</td>
<td>15 mg/kg/day</td>
</tr>
<tr>
<td>capreomycin</td>
<td>15 mg/kg IM or IV QD or 25 mg/kg PO 3 times weekly</td>
<td>1,000 mg/day</td>
</tr>
<tr>
<td>cycloserine</td>
<td>10 to 15 mg/kg PO QD or BID</td>
<td>1,000 mg/day</td>
</tr>
<tr>
<td>ethambutol</td>
<td>Follow weight-based dosing in prescribing information</td>
<td>4,000 mg/dose</td>
</tr>
<tr>
<td>ethionamide</td>
<td>10 to 20 mg/kg PO QD or BID</td>
<td>1,000 mg/day</td>
</tr>
<tr>
<td>imipenem-cilastatin*</td>
<td>1,000 mg IV BID</td>
<td>2,000 mg/day</td>
</tr>
<tr>
<td>levofloxacin</td>
<td>500 to 1,000 mg PO or IV QD</td>
<td>1,000 mg/day</td>
</tr>
<tr>
<td>linezolid</td>
<td>600 mg PO or IV QD</td>
<td>600 mg/day</td>
</tr>
<tr>
<td>meropenem*</td>
<td>2,000 mg IV BID or TID</td>
<td>6,000 mg/day</td>
</tr>
<tr>
<td>moxifloxacin</td>
<td>400 mg PO or IV QD</td>
<td>400 mg/day</td>
</tr>
<tr>
<td>para-aminosalicylic acid</td>
<td>8 to 12 g PO BID or TID</td>
<td>12 g/day</td>
</tr>
<tr>
<td>pyrazinamide</td>
<td>Follow weight-based dosing in prescribing information</td>
<td>4,000 mg/dose</td>
</tr>
<tr>
<td>Streptomycin</td>
<td>15 mg/kg IM or IV QD or 25 mg/kg PO 3 times weekly</td>
<td>20 mg/kg/day</td>
</tr>
<tr>
<td>pretomanid</td>
<td>200 mg PO QD for 26 weeks.</td>
<td>200 mg/day</td>
</tr>
</tbody>
</table>
### Drug Name | Dosing Regimen | Dose Limit/ Maximum Dose
--- | --- | ---
linezolid | 1,200 mg PO QD | 1,200 mg/day

*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.*

*Amoxicillin-clavulanic acid should be coadministered with every dose of imipenem-cilastatin or meropenem but is not counted as a separate agent and should not be used as a separate agent.*

#### Appendix C: Contraindications/Boxed Warnings
- Contraindication(s): none reported
- Boxed warning(s): increased mortality, QT prolongation

#### Appendix D: General Information

**For MDR-TB:**
- Sirturo should only be used in combination with at least 3 other drugs to which the patient’s MDR-TB isolate has been shown to be susceptible *in vitro*. If *in vitro* testing results are unavailable, Sirturo treatment may be initiated in combination with at least 4 other drugs to which the patient’s MDR-TB isolate is likely susceptible.
- Sirturo was approved under accelerated approval based on time to sputum culture conversion. Continued approval for its indication may be contingent upon verification and description of clinical benefit in confirmatory trials.
- Laboratory confirmation of multi-drug resistant TB must show TB with an isolate showing genotypic or phenotypic resistance to isoniazid and rifampin.

**For MDR-TB or XDR-TB with pretomanid:**
- Pretomanid should only be used in combination with Sirturo and linezolid.
- Dosing of the combination regimen of pretomanid, Sirturo, and linezolid can be extended beyond 26 weeks if necessary, to a maximum of 9 months, in patients with delayed culture conversion.
  - Delayed culture conversion: two consecutive negative sputum cultures following an initial positive culture.
- Laboratory confirmation of multi-drug resistant TB must show TB with an isolate showing genotypic or phenotypic resistance to isoniazid and rifampin.
- Laboratory confirmation of extensively drug resistant TB must show TB with an isolate showing genotypic or phenotypic resistance to isoniazid, rifampin, fluoroquinolones, as well as second-line injectable agents such as aminoglycosides or capreomycin.
- Linezolid starting dose of 1,200 mg daily for 26 weeks may be managed as follows:
  - Adjusted to 600 mg daily and further reduced to 300 mg daily as necessary for adverse reactions of myelosuppression, peripheral neuropathy, and optic neuropathy.
  - Doses of the regimen missed for safety reasons can be made up at the end of treatment; doses of linezolid alone missed due to adverse reactions should not be made up.
V. Dosage and Administration

<table>
<thead>
<tr>
<th>Indication</th>
<th>Dosing Regimen</th>
<th>Maximum Dose</th>
</tr>
</thead>
<tbody>
<tr>
<td>MDR-TB</td>
<td>Weight ≥ 30 kg: 400 mg PO QD for the first 2 weeks, followed by 200 mg PO three times per week (with at least 48 hours between doses) for 22 weeks (total duration of 24 weeks). Weight 15 to 29 kg: 200 mg PO QD for the first 2 weeks, followed by 100 mg PO three times per week (with at least 48 hours between doses) for 22 weeks (total duration of 24 weeks). Situro should be administered by directly observed therapy (DOT)</td>
<td>Weight ≥ 30 kg: 400 mg/dose Weight 15 to 29 kg: 200 mg/dose</td>
</tr>
<tr>
<td>MDR-TB or XDR-TB with pretomanid</td>
<td>Administer in combination with pretomanid and linezolid in a directly observed therapy (DOT) setting. • Sirturo: 400 mg PO QD for the first 2 weeks, followed by 200 mg PO three times per week (with at least 48 hours between doses) for 24 weeks (total duration of 26 weeks). • Pretomanid: 200 mg PO QD for 26 weeks. • Linezolid: 1,200 mg PO QD for 26 weeks. Patients may continue treatment with Sirturo and pretomanid without linezolid if the patient has previously received a total daily dose of linezolid 1,200 mg for at least 4 weeks.</td>
<td>400 mg/dose</td>
</tr>
</tbody>
</table>

VI. Product Availability

Tablet: 20 mg, 100 mg

VII. References


<table>
<thead>
<tr>
<th>Reviews, Revisions, and Approvals</th>
<th>Date</th>
<th>P&amp;T Approval Date</th>
</tr>
</thead>
<tbody>
<tr>
<td>Policy created: adapted from previously approved policy HIM.PA.18 (to be retired); added HIM and Medicaid lines of business; added additional therapy options to Appendix B (alphabetized table) based on 2019 WHO guidelines and commercially available in the US (ethambutol, imipenem-cilastatin, linezolid, meropenem); updated FDA-approved age limit to 12 years of age and older; references reviewed and updated.</td>
<td>08.05.19</td>
<td>11.19</td>
</tr>
<tr>
<td>Criteria added for treatment of multi-drug resistant and extensively drug resistant TB with pretomanid; Added general information regarding all oral combination regimen of pretomanid, bedaquiline, and linezolid based on FDA briefing document; references reviewed and updated.</td>
<td>09.24.19</td>
<td>02.20</td>
</tr>
<tr>
<td>RT4: updated for pediatric extension from 12 years old or 30 kg to 5 years of age or 15 kg for MDR-TB without Pretomanid per revised prescribing information.</td>
<td>06.02.20</td>
<td></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.
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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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