Clinical Policy: Linezolid (Zyvox)
Reference Number: CP.PMN.27
Effective Date: 09.01.06
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
Linezolid (Zyvox®) is an oxazolidinone-class antibacterial agent.

FDA Approved Indication(s)
Zyvox is indicated in adults and children for the treatment of the following infections caused by susceptible gram-positive bacteria:

- Nosocomial pneumonia caused by *Staphylococcus aureus* (methicillin-susceptible and -resistant isolates) or *Streptococcus pneumoniae*
- Community-acquired pneumonia caused by *Streptococcus pneumoniae*, including cases with concurrent bacteremia, or *Staphylococcus aureus* (methicillin-susceptible isolates only)
- Complicated skin and skin structure infections, including diabetic foot infections, without concomitant osteomyelitis caused by *Staphylococcus aureus* (methicillin-susceptible and -resistant isolates), *Streptococcus pyogenes*, or *Streptococcus agalactiae*. Zyvox has not been studied in the treatment of decubitus ulcers
- Uncomplicated skin and skin structure infections caused by *Staphylococcus aureus* (methicillin-susceptible isolates only) or *Streptococcus pyogenes*
- Vancomycin-resistant *Enterococcus faecium* infections, including cases with concurrent bacteremia

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 Zyvox is medically necessary when the following criteria are met:

I. Initial Approval Criteria
   A. All FDA-Approved Indications (must meet all):
      1. Diagnosis is an FDA-approved indication;
      2. Member meets one of the following (a or b):
         a. Request is for continuation of therapy initiated in an acute care hospital from which member was discharged;
         b. Both of the following (i and ii):
            i. Culture and sensitivity (C&S) report for the current infection shows isolated pathogen is a gram-positive bacteria susceptible to linezolid, unless provider submits documentation that obtaining a C&S report is not feasible;
            ii. Member meets one of the following (a, b, or c):
a) Failure of ≥ 2 formulary antibiotics to which the isolated pathogen is susceptible (if available) per C&S report, unless all are contraindicated or clinically significant adverse effects are experienced;

b) C&S report shows resistance or lack of susceptibility of the isolated pathogen to all formulary antibiotics FDA-approved for member’s diagnosis;

c) If provider documents that obtaining a C&S report is not feasible: Failure of ≥ 2 formulary antibiotics indicated for member’s diagnosis (if available), unless all are contraindicated or clinically significant adverse effects are experienced;

3. Dose does not exceed 1,200 mg (2 tablets, 2 vials, or 60 mL suspension) per day.

Approval duration:
HIM/Medicaid – Duration of request or up to 28 days of total treatment, whichever is less

B. Multi-Drug Resistant Tuberculosis with Pretomanid (off-label) (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 Sirturo® (bedaquiline) and pretomanid;
      *Prior authorization may be required for pretomanid and Sirturo.
   5. Documented resistance to fluoroquinolones, unless contraindicated or clinically significant adverse effects are experienced;
   6. Dose does not exceed 1,200 mg (2 tablets) per day.

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. All FDA-Approved Indications (must meet all):
      1. Member meets one of the following (a or b):
         a. Currently receiving medication via Centene benefit or member has previously met initial approval criteria;
         b. Request is for continuation of therapy initiated in an acute care hospital from which member was discharged;
      2. Member is responding positively to therapy;
      3. Member has not received ≥ 28 days of therapy for current infection;
      4. If request is for a dose increase, new dose does not exceed 1,200 mg (2 tablets, 2 vials, or 60 mL suspension) per day.

Approval duration:
HIM/Medicaid – Up to 28 days of total treatment
B. Multi-Drug Resistant Tuberculosis with Pretomanid (off-label) (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 in combination with Sirturo;
   4. If request is for a dose increase, new dose does not exceed 1,200 mg (2 tablets) daily.

   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 28 days (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
   C&S: culture and sensitivity
   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>pretomanid</td>
<td>200 mg PO QD for 26 weeks.</td>
<td>200 mg/day</td>
</tr>
<tr>
<td>Sirturo®</td>
<td>400 mg PO QD for the first 2 weeks, followed by 200 mg PO three times per week for remaining 24 weeks.</td>
<td>400 mg/day</td>
</tr>
</tbody>
</table>

   Therapeutic alternatives include formulary antibiotics that are indicated for member’s diagnosis and have sufficient activity against the offending pathogen at the site of the infection.

   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
• Contraindication(s):
  o Known hypersensitivity to linezolid or any of the other product components
  o Patients taking any monoamine oxidase inhibitors (MAOI) within two weeks of taking an MAOI
• Boxed warnings(s): none reported

**Appendix D: General Information**
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.
  o 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:
  o Adjusted to 600 mg daily and further reduced to 300 mg daily as necessary for adverse reactions of myelosuppression, peripheral neuropathy, and optic neuropathy.
  o Doses of the regiment 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 or XDR-TB with pretomanid</td>
<td>Linezolid to be administered in combination with Sirturo and pretomanid in a directly observed therapy (DOT) setting.</td>
<td>1,200 mg/day</td>
</tr>
<tr>
<td></td>
<td>• Take 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).</td>
<td></td>
</tr>
<tr>
<td></td>
<td>• Take pretomanid 200 mg PO QD for 26 weeks.</td>
<td></td>
</tr>
<tr>
<td></td>
<td>• Take linezolid 600 mg PO QD for 26 weeks.</td>
<td></td>
</tr>
</tbody>
</table>
### Indication | Dosing Regimen | Maximum Dose |
---|---|---|
**Pediatrics** (birth – age 11 years) | | |
**Adults and Adolescents** (age ≥ 12 years) | | |
#### Nosocomial pneumonia
10 mg/kg IV or PO every 8 hours | 600 mg IV or PO every 12 hours | 10 to 14 days |
#### Community-acquired pneumonia, including concurrent bacteremia
10 mg/kg IV or PO every 8 hours | 600 mg IV or PO every 12 hours | 14 to 28 days |
#### Complicated skin and skin structure infections
10 mg/kg IV or PO every 8 hours | 600 mg IV or PO every 12 hours | 10 to 14 days |
#### Vancomycin-resistant *Enterococcus faecium* infections, including concurrent bacteremia
10 mg/kg IV or PO every 8 hours | 600 mg IV or PO every 12 hours | 14 to 28 days |
#### Uncomplicated skin and skin structure infections
Age < 5 years: 10 mg/kg PO every 8 hours | Adults: 400 mg PO every 12 hours | 10 to 14 days |
Age 5 – 11 years: 10 mg/kg PO every 12 hours | Adolescents: 600 mg PO every 12 hours | |

### VI. Product Availability
- Injection: 200 mg, 400 mg, 600 mg
- Tablets: 600 mg
- Oral suspension: 100 mg/5 mL

### VII. References

Coding Implications
Codes referenced in this clinical policy are for informational purposes only. Inclusion or exclusion of any codes does not guarantee coverage. Providers should reference the most up-to-date sources of professional coding guidance prior to the submission of claims for reimbursement of covered services.

<table>
<thead>
<tr>
<th>HCPCS Codes</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>J2020</td>
<td>Injection, linezolid, 200 mg</td>
</tr>
</tbody>
</table>

Reviews, Revisions, and Approvals

<table>
<thead>
<tr>
<th>Removed requirement related to documentation of an FDA approved indication. Added prescriber specialty. Added that culture and sensitivity report must show pathogen susceptibility to linezolid and be dated within the last 7 days. Added requirement related to trial and failure of formulary antibiotics to which pathogen is susceptible, unless contraindicated to such therapies, or culture and sensitivity report shows resistance of pathogen to formulary antibiotics. Updated continuation criteria. Updated references. Changed guideline to new format.</th>
<th>Date</th>
<th>P&amp;T Approval Date</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>11.16</td>
<td>11.16</td>
</tr>
</tbody>
</table>

Clinical changes made to criteria:
Modified criteria to allow for cases in which obtaining C&S report is not feasible per documentation from the provider
Removed language specifying “Isolated pathogen is VRE” since VRE is gram-positive and policy covers gram positive bacteria
Added max dose requirement in initial approval criteria

Non-clinical changes made:
Converted to new template
Updated policy name to reflect linezolid tablets since the oral suspension is on the formulary and does not require a PA

<table>
<thead>
<tr>
<th>Date</th>
<th>P&amp;T Approval Date</th>
</tr>
</thead>
<tbody>
<tr>
<td>01.17</td>
<td></td>
</tr>
</tbody>
</table>
### Reviews, Revisions, and Approvals

<table>
<thead>
<tr>
<th>Updated references</th>
<th>Date</th>
<th>P&amp;T Approval Date</th>
</tr>
</thead>
<tbody>
<tr>
<td>2Q 2018 annual review: no significant changes; safety updated per safety guidance endorsed by Centene Medical Affairs; references reviewed and updated.</td>
<td>03.06.18</td>
<td>05.18</td>
</tr>
<tr>
<td>1Q 2019 annual review: added criterion line for diagnosis to be an FDA-approved indication; removed 7 day requirement for C&amp;S report and replaced it with requirement that C&amp;S report is for the current infection; clarified that pathogen susceptibility to antibiotics be demonstrated via C&amp;S report; added ‘lack of susceptibility’ as an alternative to demonstrating resistance on C&amp;S; removed criterion allowing member to meet criteria if formulary antibiotics are not indicated for member’s diagnosis, since this is incorporated into other existing criteria already; added criterion to allow member to continue treatment if it was started in an acute care hospital and member was discharged; references reviewed and updated.</td>
<td>10.30.18</td>
<td>02.19</td>
</tr>
<tr>
<td>1Q 2020 annual review: 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; removed that linezolid should be prescribed by or in consultation with an ID specialist; references reviewed and updated; revised HIM line of business.</td>
<td>09.19.19</td>
<td>02.20</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.

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.

This clinical policy is the property of the Health Plan. Unauthorized copying, use, and distribution of this clinical policy or any information contained herein are strictly prohibited. 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.