PRIOR AUTHORIZATION CRITERIA

BRAND NAME (generic)

ZYVOX (linezolid)

Status: CVS Caremark® Criteria Type: Initial Prior Authorization

POLICY

FDA-APPROVED INDICATIONS

Nosocomial Pneumonia

Zyvox is indicated for the treatment of nosocomial pneumonia caused by *Staphylococcus aureus* (methicillin-susceptible and -resistant isolates) or *Streptococcus pneumoniae*.

Community-acquired Pneumonia

Zyvox is indicated for the treatment of 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

Zyvox is indicated for the treatment of 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

Zyvox is indicated for the treatment of uncomplicated skin and skin structure infections caused by *Staphylococcus aureus* (methicillin susceptible isolates only) or *Streptococcus pyogenes*.

Vancomycin-resistant Enterococcus faecium Infections

Zyvox is indicated for the treatment of vancomycin-resistant *Enterococcus faecium* infections, including cases with concurrent bacteremia.

Limitations of Use

- Zyvox is not indicated for the treatment of Gram-negative infections. It is critical that specific Gram-negative therapy be initiated immediately if a concomitant Gram-negative pathogen is documented or suspected.
- The safety and efficacy of Zyvox formulations given for longer than 28 days have not been evaluated in controlled clinical trials.

<u>Usage</u>

To reduce the development of drug-resistant bacteria and maintain the effectiveness of Zyvox and other antibacterial drugs, Zyvox should be used only to treat infections that are proven or strongly suspected to be caused by susceptible bacteria. When culture and susceptibility information are available, they should be considered in selecting or modifying antibacterial therapy. In the absence of such data, local epidemiology and susceptibility patterns may contribute to the empiric selection of therapy.

Off Label Uses

Combination regimen with Pretomanid and linezolid for the treatment of adults with pulmonary tuberculosis (TB) resistant to isoniazid, rifamycins, a fluoroquinolone and a second line injectable antibacterial drug OR adults with pulmonary TB resistant to isoniazid and rifampin, who are treatment-intolerant or nonresponsive to standard therapy.^{2, 9,10}

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COVERAGE CRITERIA

Community-Acquired Pneumonia, Complicated Skin and Skin Structure Infections, Nosocomial Pneumonia, Uncomplicated Skin and Skin Structure Infections

Authorization may be granted for the requested drug when ALL of the following criteria are met:

- The patient has ANY of the following:
 - An infection caused by vancomycin-resistant Enterococcus faecium including cases with concurrent bacteremia
 - A nosocomial (institution-acquired) 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)
 - A complicated skin and skin structure infection including diabetic foot infections, without concomitant osteomyelitis, caused by Staphylococcus aureus (methicillin-susceptible and -resistant isolates)
 Streptococcus pyogenes, or Streptococcus agalactiae
 - An uncomplicated skin and skin structure infection caused by Staphylococcus aureus (methicillin-susceptible isolates only) or Streptococcus pyogenes
- The infection is proven or strongly suspected to be caused by susceptible bacteria
- The patient meets ONE of the following:
 - The patient has experienced an inadequate treatment response, intolerance, or contraindication to alternative therapies
 - o The bacteria are NOT susceptible to any other antibiotics
- The patient will use the requested drug orally or intravenously

Non-Tuberculosis (TB) Bacterial Infection

Authorization may be granted when the requested drug is being prescribed for a NON-Tuberculous (TB) bacterial infection when ALL of the following criteria are met:

- The patient is being converted from intravenous (IV) linezolid (Zyvox)
- The requested drug is being prescribed or directed by an Infectious Disease specialist
- The patient will use the requested drug orally or intravenously

Tuberculosis (TB)

Authorization may be granted when the requested drug is being prescribed for pulmonary tuberculosis (TB) resistant to isoniazid, rifamycins, a fluoroquinolone and a second line injectable antibacterial drug OR TB resistant to isoniazid and rifampin, that is treatment-intolerant or nonresponsive to standard therapy when ALL of the following criteria are met:

- The requested drug is being prescribed as part of a combination regimen with Pretomanid and Sirturo (bedaquiline)
- The patient will use the requested drug orally or intravenously

DURATION OF APPROVAL (DOA)

- 263-A:
 - Non-Tuberculosis bacterial infection DOA: 28 days
 - o Pulmonary tuberculosis (TB) DOA: 12 months

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- 4. Diagnosis and Treatment of Adults with Community-Acquired Pneumonia. An Official Clinical Practice Guideline of the American Thoracic Society and Infectious Diseases Society of America. American Journal of Respiratory and Critical Care Medicine, Volume 200, Issue 7, 1 October 2019, Pages e45-e67.

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- 8. Pretomanid [package insert]. Hyderabad, India: Mylan Laboratories Limited for The Global Alliance for TB Drug Development (TB Alliance); April 2020.
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