

Clinical Policy: Ozenoxacin (Xepi)

Reference Number: CP.PMN.119

Effective Date: 06.01.18

Last Review Date: 05.20

Line of Business: Commercial, HIM, Medicaid

[Revision Log](#)

See [Important Reminder](#) at the end of this policy for important regulatory and legal information.

Description

Ozenoxacin (Xepi[™]) is a quinolone antimicrobial.

FDA Approved Indication(s)

Xepi is indicated for the topical treatment of impetigo due to *Staphylococcus aureus* or *Streptococcus pyogenes* in adult and pediatric patients 2 months of age and older.

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 Xepi is **medically necessary** when the following criteria are met:

I. Initial Approval Criteria

A. Impetigo (must meet all):

1. Diagnosis of impetigo;
2. Age \geq 2 months;
3. Failure of mupirocin 2% ointment or cream at up to maximally indicated doses, unless contraindicated or clinically significant adverse effects are experienced;
4. Dose does not exceed twice daily topical application for five days.

Approval duration: 1 month (1 tube)

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. Impetigo (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 twice daily topical application for five days.

Approval duration: 1 month (1 tube)

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 1 month (1 tube) (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.

| Drug Name | Dosing Regimen | Dose Limit/ Maximum Dose |
|--|--|---------------------------------------|
| mupirocin (Bactroban [®]) 2% cream, ointment | Apply small amount to affected area (up to 10 cm in length or 100 cm ² in area) TID x 10 days | TID application as outlined x 10 days |

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

| Indication | Dosing Regimen | Maximum Dose |
|------------|--|--------------------------------------|
| Impetigo | BID topical application (thin layer) for five days (for up to 100 cm ² in patients ≥ 12 years or 2% of the total body surface area and not exceeding 100 cm ² if age < 12 years. | BID application as outlined x 5 days |

VI. Product Availability

Tube containing 1% cream: 30 g

VII. References

1. Xepi Prescribing Information. Fairfield, NJ: Medimetrick's Pharmaceuticals, Inc.; January 2019. Available at: <https://dailymed.nlm.nih.gov/dailymed/drugInfo.cfm?setid=3361dd6c-4b03-4c42-9b95-4ecd43c34294>. Accessed February 6, 2020.
2. Bactroban Cream Prescribing Information. Research Triangle Park, NC: GlaxoSmithKline; March 2017. Available at https://www.accessdata.fda.gov/drugsatfda_docs/label/2017/050746s021lbl.pdf. Accessed February 6, 2020.
3. Bactroban Ointment Prescribing Information. Research Triangle Park, NC: GalxoSmithKline; March 2017. Available at: https://www.accessdata.fda.gov/drugsatfda_docs/label/2017/050703s017lbl.pdf. Accessed February 6, 2020.
4. Stevens DL, Bisno AL, Chambers HF, et al. Practice guidelines for the diagnosis and management of skin and soft tissue infections: 2014 Update by the Infectious Diseases Society of America. *Clinical Infectious Diseases* 2014;59(2):e10–52
5. Koning S, van der Sande R, Verhagen AP, et al. Interventions for impetigo (review). *Cochrane Database of Systematic Reviews*. 2012, Issue 1. Art. No.: CD003261.
6. Hartman-Adams H, Banvard C, Juckett G. Impetigo: diagnosis and treatment. *Am Fam Physician*. 2014;90(4):229-235.

| Reviews, Revisions, and Approvals | Date | P&T Approval Date |
|---|----------|-------------------|
| Policy created. | 01.30.18 | 05.18 |
| 2Q 2019 annual review: no significant changes; product availability updated; references reviewed and updated. | 02.24.19 | 05.19 |
| 2Q 2020 annual review: no significant changes; references reviewed and updated. | 02.06.20 | 05.20 |

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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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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