

Clinical Policy: Cyclosporine (Restasis)

Reference Number: CP.PMN.48

Effective Date: 05.01.12

Last Review Date: 05.18

Line of Business: Commercial, HIM, Medicaid

[Revision Log](#)

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

Description

Cyclosporine ophthalmic emulsion (Restasis[®]) is a topical calcineurin inhibitor immunosuppressant.

FDA Approved Indication(s)

Restasis is indicated to increase tear production in patients whose tear production is presumed to be suppressed due to ocular inflammation associated with keratoconjunctivitis sicca. Increased tear production was not seen in patients currently taking topical anti-inflammatory drugs or using punctal plugs.

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

I. Initial Approval Criteria

A. Keratoconjunctivitis Sicca (must meet all):

1. Diagnosis of keratoconjunctivitis sicca with suppressed tear production due to ocular inflammation;
2. Age \geq 16 years;
3. Failure of artificial tears at up to maximally indicated doses unless contraindicated or clinically significant adverse effects are experienced;
4. Failure of ophthalmic anti-inflammatory agents at up to maximally indicated doses unless contraindicated or clinically significant adverse effects are experienced;
5. Request does not exceed 60 vials per 30 days.

Approval duration:

HIM – 6 months

Commercial/Medicaid – 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. Keratoconjunctivitis Sicca (must meet all):

1. Currently receiving medication via Centene benefit or member has previously met initial approval criteria;
2. Documentation of positive response to therapy;
3. If request is for a dose increase, request does not exceed 60 vials per 30 days.

Approval duration:

HIM – 12 months

Commercial/Medicaid – 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 12 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.

Drug Name	Dosing Regimen	Dose Limit/ Maximum Dose
artificial tears (e.g., Visine dry eye relief)	1 to 2 drops in affected eye(s) BID or QID	various
ophthalmic anti-inflammatory agents (e.g., loteprednol etabonate)	1 to 2 drops in each eye BID to QID for up to 2 weeks	various

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: General Information

- Artificial tears are the standard therapy for all severity of dry eyes.

- Restasis is likely to be given in conjunction with artificial tears.
- Increased tear production was not seen in patients currently taking topical anti-inflammatory drugs or using punctal plugs.
- Emulsion from one individual single-use vial is to be used immediately after opening for administration to one or both eyes, and the remaining contents should be discarded immediately after administration.

V. Dosage and Administration

Indication	Dosing Regimen	Maximum Dose
Moderate to severe keratoconjunctivitis sicca	1 drop BID in each eye approximately 12 hours apart	2 drops/day in each eye; 60 vials/30 days

VI. Product Availability

Single use vial: 0.05%, 0.4 mL each of 30 vials/tray and 60 vials/tray
MultiDose bottle: 0.05%, 5.5 mL total

VII. References

1. Restasis Prescribing Information. Irvine, CA: Allergan, Inc.; July 2017. Available at: <https://www.restasis.com/>. Accessed January 2018.
2. The International Dry Eye Workshop. Ocul Surf. 2007; 5(2):65-204.
3. American Academy of Ophthalmology Retina Panel. Preferred Practice Pattern[®] Guidelines. Dry Eye Syndrome. San Francisco, CA: American Academy of Ophthalmology; October, 2013. Available at: www.aao.org/ppp. Accessed January 2018.
4. Restasis. Micromedex[®] Healthcare Series [Internet database]. Greenwood Village, Colo: Thomson Healthcare. Updated periodically. Accessed January 2018.

Reviews, Revisions, and Approvals	Date	P&T Approval Date
Medicaid: Removed “for a period of 3 months” related to ophthalmic anti-inflammatory use in the Criteria for Approval item D.	05.13	05.13
Medicaid: Updated references.	05.14	05.14
Medicaid: Updated references.	04.15	04.15
Medicaid: Converted to new template; Removed requirement of “failure of environmental stress reduction (use of humidifiers; consciously attempting to increase the frequency of blinking; washing the lids with a mild soap solution; and application of warm compresses)”; Updated references.	02.16	05.16
Medicaid: No clinical changes to criteria: <ul style="list-style-type: none"> • Converted to new template • Removed age criteria as age is not an absolute contraindication per FDA labeling Updated references	01.17	05.17

Reviews, Revisions, and Approvals	Date	P&T Approval Date
2Q 2018 annual review: combined Medicaid, HIM, and commercial lines of business criteria; commercial: removed ophthalmologist or optometrist prescriber requirement; expanded requirement of any OTC wetting agent to artificial tears and anti-inflammatory agent; Medicaid: expanded approval duration from 6 months (initial) and 12 months (continued) to length of benefit	02.02.18	05.18

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.

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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