

PRIOR AUTHORIZATION POLICY

POLICY: Ophthalmology – Dry Eye Disease – Eysuvis Prior Authorization Policy

Eysuvis® (loteprednol etabonate 0.25% ophthalmic suspension –

Kala)

REVIEW DATE: 12/06/2023

INSTRUCTIONS FOR USE

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CIGNA NATIONAL FORMULARY COVERAGE:

OVERVIEW

Eysuvis, an ophthalmic corticosteroid, is indicated for the **short-term (up to 2** weeks) treatment of the signs and symptoms of dry eye disease.¹

Guidelines

Eysuvis is not addressed in guidelines. The American Academy of Ophthalmology published a Preferred Practice Pattern® (2018) for the treatment of dry eye syndrome.³ For mild dry eyes, education and environmental modifications, artificial tear solutions, and eyelid therapy (warm compresses and eyelid scrubs) are listed as some of the treatment options. The guidelines note commercially available loteprednol etabonate 0.5% was used in a prospective, randomized study for a 2-week period. The study found a favorable effect in patients' dry eye symptoms and conjunctival hyperemia findings.

POLICY STATEMENT

Prior Authorization is recommended for prescription benefit coverage of Eysuvis. All approvals are provided for the duration noted below. In cases where the approval is authorized in months, 1 month is equal to 30 days.

 Eysuvis® (loteprednol etabonate 0.25% ophthalmic suspension – Kala)

is(are) covered as medically necessary when the following criteria is(are) met for FDA-approved indication(s) or other uses with supportive evidence (if applicable):

FDA-Approved Indication

1. Dry Eye Disease (Short-Term Treatment). Approve for 1 month if the patient has tried artificial tears.

CONDITIONS NOT COVERED

Eysuvis® (loteprednol etabonate 0.25% ophthalmic suspension – Kala) is(are) considered experimental, investigational or unproven for ANY other use(s).

REFERENCES

- 1. Eysuvis[™] ophthalmic suspension [prescribing information]. Watertown, MA: Kala; July 2022.
- 2. Korenfeld M, Nichols KK, Goldberg D, et al. Safety of KPI-121 ophthalmic suspension 0.25% in patients with dry eye disease: A pooled analysis of 4 multicenter, randomized, vehicle-controlled studies. *Cornea*. 2021 May 1;40(5):564-570.
- 3. Akpek EK, Amescua G, Farid M, et al. Dry Eye Syndrome Preferred Practice Pattern®. *Ophthalmology*. 2019 Jan;126(1):P286-P334.

HISTORY

Type of Revision	Summary of Changes	Review Date
Annual Revision	No criteria changes.	12/07/2022
Annual Revision	No criteria changes.	12/06/2023

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