PHARMACY POLICY STATEMENT
Indiana Medicaid

DRUG NAME	Yutiq (fluocinolone acetonide)
BENEFIT TYPE	Medical
STATUS	Prior Authorization Required

Yutiq is a 0.18 mg fluocinolone acetonide intravitreal implant that was approved by the FDA in 2018. It is indicated for the treatment of chronic non-infectious uveitis affecting the posterior segment of the eye and lasts 36 months.
Uveitis is an inflammation of the uvea (middle layer of the eye). It can be infectious or non-infectious. Non-infectious uveitis (NIU) is often associated with inflammatory conditions such as rheumatoid arthritis. If the anterior segment of the uvea is affected, it can be treated with topical glucocorticoids. If resistant or affecting the intermediate or posterior segments, more invasive or systemic treatment is needed.

Yutiq (fluocinolone acetonide) will be considered for coverage when the following criteria are met:

Uveitis

For **initial** authorization:
1. Member is at least 18 years of age; AND
2. Medication must be prescribed by or in consultation with an ophthalmologist; AND
3. Member has a diagnosis of chronic (1 year or more) non-infectious uveitis affecting the posterior segment of the eye; AND
4. Member has tried and failed at least one of the following for at least 3 months:
   a) Systemic corticosteroid (e.g., prednisone)
   b) Non-biologic immunosuppressive (e.g., mycophenolate mofetil, methotrexate, cyclosporine, tacrolimus); AND
5. Member does not have any active or suspected infections in or around the eye.
6. **Dosage allowed/Quantity limit:** One implant (0.18 mg) per eye
   Limit: 2 implants (1 per eye) per 36 months

If all the above requirements are met, the medication will be approved for 3 months.

For **reauthorization**:
1. Chart notes must show improved or stabilized visual acuity following treatment and/or an improved vitreous haze score; AND

CareSource considers Yutiq (fluocinolone