INDICATIONS AND USAGE

**YUTIQ™** (fluocinolone acetonide intravitreal implant) 0.18 mg is indicated for the treatment of chronic non-infectious uveitis affecting the posterior segment of the eye.

Please see Important Safety Information throughout and on page 11 and full Prescribing Information in pocket.
YUTIQ is designed to deliver a sustained release of fluocinolone for patients with chronic noninfectious posterior uveitis for up to 36 months¹

- Each non-bioerodible intravitreal implant contains 0.18 mg of fluocinolone acetonide released at an initial rate of 0.25 mcg/day over 36 months
- The implant measures 3.5 mm x 0.37 mm
- YUTIQ (fluocinolone acetonide intravitreal injection) 0.18 mg is supplied in a sterile single-dose preloaded applicator with a 25-gauge needle, packaged in a sealed sterile pouch

Utilizes micro-implant Durasept™ Technology from EyePoint™²

IMPORTANT SAFETY INFORMATION
Contraindications
Ocular or Periocular Infections: YUTIQ is contraindicated in patients with active or suspected ocular or periocular infections including most viral disease of the cornea and conjunctiva including active epithelial herpes simplex keratitis (dendritic keratitis), vaccinia, varicella, mycobacterial infections and fungal diseases.

Please see Important Safety Information throughout and on page 11 and full Prescribing Information in pocket.
Preparing the patient for the intravitreal injection of YUTIQ

- The intravitreal injection procedure should be carried out under aseptic conditions, which includes sterile materials: gloves, drape, caliper, and eyelid speculum (or equivalent). Adequate anesthesia and a broad-spectrum microbicide should be given prior to the injection.

STEP 1

- Just prior to injection, administer topical and/or subconjunctival anesthesia at the injection site (inferotemporal quadrant recommended)

Contraindications (cont’d)

Hypersensitivity: YUTIQ is contraindicated in patients with known hypersensitivity to any components of this product.
YUTIQ™ injection procedure

STEP 2

- Administer 2 to 3 drops of a broad-spectrum microbicide into the lower fornix
- The lids may be scrubbed with cotton-tipped applicators soaked with a broad-spectrum microbicide
- Place a sterile lid speculum
- Have the patient look up and apply additional microbicide solution to the injection site
- Allow 30 to 60 seconds for the topical antiseptic to dry prior to injection of YUTIQ (fluocinolone acetonide intravitreal implant) 0.18 mg

STEP 3

- Optimal placement of YUTIQ is inferior to the optic disc and posterior to the equator of the eye
- Measure 4 mm inferotemporal from the limbus with the aid of calipers for point of entry into the sclera

IMPORTANT SAFETY INFORMATION (cont’d)

Warnings and Precautions

Intravitreal Injection-related Effects: Intravitreal injections, including those with YUTIQ, have been associated with endophthalmitis, eye inflammation, increased or decreased intraocular pressure, and choroidal or retinal detachments. Hypotony has been observed within 24 hours of injection and has resolved within 2 weeks. Patients should be monitored following the intravitreal injection.

Please see Important Safety Information throughout and on page 11 and full Prescribing Information in pocket.
STEP 3

• Optimal placement of YUTIQ is inferior to the optic disc and posterior to the equator of the eye
• Measure 4 mm inferotemporal from the limbus with the aid of calipers for point of entry into the sclera

STEP 2

• Administer 2 to 3 drops of a broad-spectrum microbicide into the lower fornix
• The lids may be scrubbed with cotton-tipped applicators soaked with a broad-spectrum microbicide
• Place a sterile lid speculum
• Have the patient look up and apply additional microbicide solution to the injection site
• Allow 30 to 60 seconds for the topical antiseptic to dry prior to injection of YUTIQ (fluocinolone acetonide intravitreal implant) 0.18 mg

Warnings and Precautions (cont’d)
Steroid-related Effects: Use of corticosteroids including YUTIQ may produce posterior subcapsular cataracts, increased intraocular pressure and glaucoma. Use of corticosteroids may enhance the establishment of secondary ocular infections due to bacteria, fungi, or viruses. Corticosteroids are not recommended to be used in patients with a history of ocular herpes simplex because of the potential for reactivation of the viral infection.
STEP 6

• Remove the black plunger stop from the plunger

STEP 7

• Carefully remove the protective cap from the needle and inspect the needle tip to ensure it is not bent

IMPORTANT SAFETY INFORMATION (cont’d)

Warnings and Precautions (cont’d)

Risk of Implant Migration: Patients in whom the posterior capsule of the lens is absent or has a tear are at risk of implant migration into the anterior chamber.

Please see Important Safety Information throughout and on page 11 and full Prescribing Information in pocket.
STEP 8

• Remove the trombone wire from the distal end of the needle. Prior to injection, keep the applicator tip above the horizontal plane to ensure that the YUTIQ (fluocinolone acetonide intravitreal implant) 0.18 mg implant does not fall out of the applicator.

STEP 9

• Gently displace the conjunctiva so that after withdrawing the needle, the conjunctival and scleral needle entry sites will not align.
• Care should be taken to avoid contact between the needle and the lid margin or lashes.

Adverse Reactions

In controlled studies, the most common adverse reactions reported were cataract development and increases in intraocular pressure.
STEP 10

- Insert the needle through the conjunctiva and sclera up to the positive stop of the applicator

STEP 11

- Depress the plunger at the back of the applicator fully to deliver the YUTIQ (fluocinolone acetonide intravitreal implant) 0.18 mg implant into the back of the eye

IMPORTANT SAFETY INFORMATION

Contraindications

Ocular or Periocular Infections: YUTIQ is contraindicated in patients with active or suspected ocular or periocular infections including most viral disease of the cornea and conjunctiva including active epithelial herpes simplex keratitis (dendritic keratitis), vaccinia, varicella, mycobacterial infections and fungal diseases.

Please see Important Safety Information throughout and on page 11 and full Prescribing Information in pocket.
STEP 11
• Depress the plunger at the back of the applicator fully to deliver the YUTIQ (fluocinolone acetonide intravitreal implant) 0.18 mg implant into the back of the eye. Please see Important Safety Information throughout and on page 11 and full Prescribing Information in pocket.

Contraindications

Ocular or Periocular Infections:
YUTIQ is contraindicated in patients with active or suspected ocular or periocular infections including most viral disease of the cornea and conjunctiva including active epithelial herpes simplex keratitis (dendritic keratitis), vaccinia, varicella, mycobacterial infections and fungal diseases.

STEP 10
• Insert the needle through the conjuctiva and sclera up to the positive stop of the applicator.

STEP 12
• Remove the YUTIQ applicator from the eye and discard in biohazard sharps container.

STEP 13
• Remove the lid speculum and perform indirect ophthalmoscopy to verify adequate central retinal artery perfusion, absence of any other complications, and to verify the placement of the implant.
• Scleral depression may enhance visualization of the implant.
• Immediate measurement of intraocular pressure (IOP) may be performed at the discretion of the ophthalmologist.

Contraindications (cont’d)

Hypersensitivity: YUTIQ is contraindicated in patients with known hypersensitivity to any components of this product.
Follow up after YUTIQ™ insertion

- Monitor patients for changes in IOP and endophthalmitis following injection. Monitoring may consist of:
  - A check for perfusion of the optic nerve head immediately after the injection
  - Tonometry within 30 minutes following injection
  - Biomicroscopy between 2 and 7 days following injection
- Patients should be instructed to report without delay any symptoms suggestive of endophthalmitis.
INDICATIONS AND USAGE
YUTIQ™ (fluocinolone acetonide intravitreal implant) 0.18 mg is indicated for the treatment of chronic non-infectious uveitis affecting the posterior segment of the eye.

IMPORTANT SAFETY INFORMATION
Contraindications
Ocular or Periocular Infections: YUTIQ is contraindicated in patients with active or suspected ocular or periocular infections including most viral disease of the cornea and conjunctiva including active epithelial herpes simplex keratitis (dendritic keratitis), vaccinia, varicella, mycobacterial infections and fungal diseases.

Hypersensitivity: YUTIQ is contraindicated in patients with known hypersensitivity to any components of this product.

Warnings and Precautions
Intravitreal Injection-related Effects: Intravitreal injections, including those with YUTIQ, have been associated with endophthalmitis, eye inflammation, increased or decreased intraocular pressure, and choroidal or retinal detachments. Hypotony has been observed within 24 hours of injection and has resolved within 2 weeks. Patients should be monitored following the intravitreal injection.

Steroid-related Effects: Use of corticosteroids including YUTIQ may produce posterior subcapsular cataracts, increased intraocular pressure and glaucoma. Use of corticosteroids may enhance the establishment of secondary ocular infections due to bacteria, fungi, or viruses. Corticosteroids are not recommended to be used in patients with a history of ocular herpes simplex because of the potential for reactivation of the viral infection.

Risk of Implant Migration: Patients in whom the posterior capsule of the lens is absent or has a tear are at risk of implant migration into the anterior chamber.

Adverse Reactions
In controlled studies, the most common adverse reactions reported were cataract development and increases in intraocular pressure.

Please see full Prescribing Information in pocket.
INDICATIONS AND USAGE
YUTIQ™ (fluocinolone acetonide intravitreal implant) 0.18 mg is indicated for the treatment of chronic non-infectious uveitis affecting the posterior segment of the eye.

IMPORTANT SAFETY INFORMATION
Contraindications
Ocular or Periocular Infections: YUTIQ is contraindicated in patients with active or suspected ocular or periocular infections including most viral disease of the cornea and conjunctiva including active epithelial herpes simplex keratitis (dendritic keratitis), vaccinia, varicella, mycobacterial infections and fungal diseases.

Hypersensitivity: YUTIQ is contraindicated in patients with known hypersensitivity to any components of this product.


Please see Important Safety Information throughout and on page 11 and full Prescribing Information in pocket.