

For diabetic macular edema (DME)

**ILUVIEN**<sup>®</sup>  
(fluocinolone acetonide  
intraocular implant) 0.19mg

**ONE INJECTION**  
Up to 3 years of continuous  
treatment for diabetic  
macular edema (DME)

**IMAGINE**

**WHAT YOU'LL SEE**

Fewer injections could help you  
embrace life's magical moments



**Ask your eye doctor if  
prescription ILUVIEN is right for you.**

**Visit [ILUVIEN.com](http://ILUVIEN.com) or call us at 1-844-445-8843**

**Please see Important Safety Information on back panel  
and accompanying full Prescribing Information.**

### **Important Safety Information**

- Do not use ILUVIEN if you have or think you might have an infection in or around the eye.
- ILUVIEN should not be used if you have advanced glaucoma.
- You should not use ILUVIEN if you are allergic to any ingredients of ILUVIEN.

## Indication and Important Safety Information

### Important Safety Information (continued)

- Injections into the vitreous in the eye are associated with a serious eye infection (endophthalmitis), eye inflammation, increased eye pressure, glaucoma, and retinal detachments. Your eye doctor should monitor you regularly after the injection.
- Use of corticosteroids including ILUVIEN may produce cataracts (ILUVIEN 82%; sham 50%), increased eye pressure (ILUVIEN 34%; sham 10%), glaucoma, and may increase secondary eye infections due to bacteria, fungi, or viruses. Let your doctor know if you have a history of herpes viral infections of the eye.
- If the posterior capsule of the lens of your eye is missing or torn the ILUVIEN implant may move to the front chamber of the eye.
- The most common side effects reported in patients with diabetic macular edema who were treated with ILUVIEN include cataracts (ILUVIEN 82%; sham 50%) and increased eye pressure (ILUVIEN 34%; sham 10%).

**Indication:** ILUVIEN® (fluocinolone acetonide intravitreal implant) 0.19 mg is indicated for the treatment of diabetic macular edema (DME) in patients who have been previously treated with a course of corticosteroids and did not have a clinically significant rise in intraocular pressure.

You are encouraged to report negative side effects of prescription drugs to the FDA. Visit [www.fda.gov/medwatch](http://www.fda.gov/medwatch), or call **1-800-FDA-1088**.

Please see accompanying full Prescribing Information.



To learn more, visit [ILUVIEN.com](http://ILUVIEN.com) by using your mobile device to scan the QR code.

**ALIMERA**  
SCIENTIFICS