MEET KELLY

42-year-old female

Digital artist; spends most of the day working on a tablet or laptop

Medical History

- **No significant** ocular history
- Has self-medicated with **multiple OTC drops over the last year** but isn't sure which ones; found them **ineffective**
- Takes a **daily antihistamine**

Chief Complaints

- Increasing **difficulty with near vision**; patient **believes she requires glasses**
- Symptoms of **eye fatigue, burning eyes,** and **fluctuating vision,** especially late in the day
- Reports a **gritty and scratchy feeling** in her eyes. Visual symptoms and dry, irritated eyes **affect her ability to perform her work**

Initial Observations

- Mild eye **redness**
- **Blinks to improve vision** during visual acuity testing

INDICATION

MIEBO™ (perfluorohexyloctane ophthalmic solution) is a semifluorinated alkane indicated for the treatment of the signs and symptoms of dry eye disease.

IMPORTANT SAFETY INFORMATION

- MIEBO should not be administered while wearing contact lenses. Contact lenses should be removed before use and for at least 30 minutes after administration of MIEBO

Please see additional Important Safety Information on next page.

Click here for full Prescribing Information for MIEBO.

Miebo™
(perfluorohexyloctane ophthalmic solution)

MEET KELLY

NEXT STEPS

Based on Kelly's history and presentation, **what assessments would you perform?**

What results would you expect to find?

What would be **your proposed treatment plan** to address Kelly's symptoms?

Discover how MIEBO could provide patients like Kelly with **relief from the signs and symptoms** of dry eye disease.

Learn more at MIEBO-ECP.COM.

IMPORTANT SAFETY INFORMATION (CONTINUED)

- Instruct patients to instill one drop of MIEBO into each eye four times daily
- The safety and efficacy in pediatric patients below the age of 18 have not been established
- The most common ocular adverse reaction was blurred vision (1% to 3% of patients reported blurred vision and conjunctival redness)

You are encouraged to report negative side effects of prescription drugs to the FDA. Visit www.fda.gov/medwatch or call 1-800-FDA-1088.

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