



# Rx STRENGTH LASTACAFT AVAILABLE OTC

FOR PATIENTS WITH ITCHING DUE TO ALLERGIC CONJUNCTIVITIS



**FAST-ACTING:** Effective at 3 Minutes LONG-LASTING: Once Daily Relief

- 2x the amount per bottle than PATADAY®\*
- Works against common allergens: pet dander, pollen, grass, and ragweed
- For ages 2+

\*60 day supply vs. PATADAY® 30 day supply

IT'S LASTACAFT® SEASON

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## **MECHANISM OF ACTION:**

H1 Histamine Receptor Antagonist Inhibitor of Histamine Release from Mast Cells

# CLINICALLY PROVEN TO RELIEVE OCULAR ALLERGY ITCH AT 3 MINUTES AND THROUGH 16 HOURS

#### FAST-ACTING: effective at 3 minutes<sup>1-3</sup>

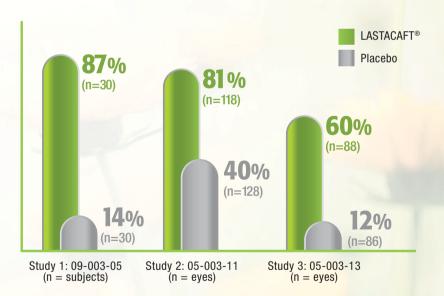
15 minutes post-dosing (3 minutes post-CAC)

#### 100 LASTACAFT® **97**% 90 Placebo (n=29)80 PERCENTAGE OF SUBJECTS REPORTING MINIMAL ITCH<sup>a</sup> **81**% 70 (n=122)**70**% 60 (n=88)50 40 30 20 **23**% **24**% **17**% 10 (n=130)(n=84)(n=29)Study 1: 09-003-05 Study 2: 05-003-11 Study 3: 05-003-13

(n = eyes)

### LONG-LASTING: effective through 16 hours<sup>1-3</sup>

16 hours post-dosing (3 minutes post-CAC)



<sup>a</sup>Minimal itch defined as an ocular itch score from 0 to < 1.

Do not use if solution changes color or becomes cloudy, if sensitive to any ingredient in this product, or to treat contact lens related irritation.

#### **Study Design and Methodology**

(n = subjects)

All clinical efficacy trials were phase 3, double-masked, randomized, vehicle-controlled studies. These 3 trials included 274 patients ≥ 10 years of age in the intent-to-treat population. The Conjunctival Allergen Challenge (CAC) model was used in all 3 studies to test the effect of LASTACAFT® at onset and duration of action. Each study included 4 visits over approximately 5 weeks, and each patient received a total of 2 doses of LASTACAFT® or placebo.<sup>4</sup>

At visit 3, the CAC was conducted 16 hours post dosing of LASTACAFT® or placebo, and the results shown were obtained 3 minutes post-CAC. At visit 4, the CAC was conducted 15 minutes post dosing of LASTACAFT® or placebo, and the results shown were obtained 3 minutes post-CAC. The main study end point was ocular itching evaluated by the patient at 3, 5, and 7 minutes post-CAC at visits 3 and 4. Ratings were made on a scale of 0 to 4 (allowing half-unit increments), where 0 = "none" and 4 = "an incapacitating itch with an irresistible urge to rub." 1-4

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