

## **Patient Consent for Latisse®**

Latisse® is a prescription treatment approved by the FDA to grow longer, thicker, and darker lashes.

Latisse® is believed to affect the growth phase of the eyelash hair cycle in 2 ways:

- 1. Increase the length of the "growth" phase.
- 2. Increase the number of hairs in the "growth" phase

The exact way in which Latisse® works is still unknown. The onset of results is gradual but usually a significant improvement is noted after 2 months of nightly use. Additional application is not known to increase the "growth" phase.

## **Contraindications**

You should not use Latisse® if you are <u>ALLERGIC</u> to one of its ingredients. Also it is important to let you physician <u>KNOW</u> if you answered "YES" to any of the following questions:

<ul> <li>Do you have a family member (blood relative) with GLAUCOMA? Yes No</li></ul>	Si	gnature Date Witness Date
<ul> <li>Do you have a family member (blood relative) with GLAUCOMA? Yes No</li></ul>	giv	
<ul> <li>Do you have a family member (blood relative) with GLAUCOMA? Yes No</li></ul>	ha	you develop a new eye condition (ex: eye trauma or infection), experience a sudden decrease in vision we eye or eye lid surgery, or develop any eye or eyelid reactions, particularly an infection, or if you come pregnant, you should immediately talk to your <b>DOCTOR</b> about whether to continue Latisse® usage.
<ul> <li>Do you have a family member (blood relative) with GLAUCOMA? Yes No</li> <li>Do you have a history of elevated INTRAOCULAR PRESSURE? Yes No</li> <li>Are you currently taking medication for eye pressure problems? Yes No</li> <li>Are you pregnant or breast feeding? Yes No</li> <li>Side effects</li> <li>Possible side effects (in approximately 4% of reported users in Latisse® clinical trial) include: <ul> <li>Eye itching</li> </ul> </li> </ul>	•	Eye irritation  Dryness or redness of the eyelids  Darkening of eyelid skin which may be reversible  Darkening of the eye color, for example, hazel, green, or blue eyes change to brown (possibly permanent). However this was not seen in clinical trials for Latisse®. Documentation of higher doses of bimatoprost ophthalmic solution at significantly higher concentrations and when used for treatment of glaucoma.
<ul> <li>Do you have a family member (blood relative) with GLAUCOMA? Yes No</li> <li>Do you have a history of elevated INTRAOCULAR PRESSURE? Yes No</li> <li>Are you currently taking medication for eye pressure problems? Yes No</li> <li>Are you pregnant or breast feeding? Yes No</li> <li>Side effects</li> </ul>		Eye redness
<ul> <li>Do you have a family member (blood relative) with GLAUCOMA? Yes No</li> <li>Do you have a history of elevated INTRAOCULAR PRESSURE? Yes No</li> <li>Are you currently taking medication for eye pressure problems? Yes No</li> <li>Are you pregnant or breast feeding? Yes No</li> </ul>		
<ul> <li>Do you have a family member (blood relative) with GLAUCOMA? Yes No</li> <li>Do you have a history of elevated INTRAOCULAR PRESSURE? Yes No</li> </ul>	•	Are you pregnant or breast feeding? Yes No
		Do you have a history of elevated INTRAOCULAR PRESSURE? Yes No