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Informed Consent for Latisse

Full Name _____Age _____ Date _____

Latisse™ is the brand name for bimatoprost . Latisse is FDA approved for the treatment of hypotrichosis of the eyelashes by making them grow longer, thicker and darker. Hypotrichosis is a medical term for short or missing lashes. Latisse is believed to affect the growth (anagen) phase of the eyelash hair cycle by increasing the length of the growth phase and increasing the number of hairs along the eyelid margin. The onset of action is gradual with most users seeing a significant improvement in the length and number of lashes by 2 months. If Latisse is discontinued the eyelashes and eyelids will return to their previous appearance over several weeks to months. The following side effects are the most frequently reported, but occur in less than 4% of users (i.e. 4 out of 100 users): Eye irritation and itching, conjunctival hyperemia or red eye, dry eye symptoms, and eyelid redness. Although rare, Latisse has the potential to permanently increase the brown pigmentation of the iris (colored part of the eyeball, inside the eye). Latisse may cause hyperpigmentation or darkening of the eyelid skin which may or may not be reversible upon discontinuation of the treatment. Latisse may lower intraocular pressure (IOP) or pressure inside the eye; however, the magnitude of this reduction is usually not a cause for concern. If you have a history of abnormal eye pressures or glaucoma you should only use Latisse under the close supervision of your ophthalmologist. Inform anyone conducting an eye pressure examination that you are using Latisse. You should inform your ophthalmologist that you are using Latisse if eye surgery is planned. Latisse is intended for use on the skin at the base of the eyelashes of the UPPER eyelids only. DO NOT APPLY to the lower eyelids as this will increase the chance of side effects such as hyperpigmentation or darkening of the eyelid skin. You should NOT use Latisse if: you are allergic or hypersensitive to bimatoprost (LumiganR) or any other ingredient in this product; are about to undergo cataract or other eye procedures, have an intraocular inflammation (uveitis), have risk factors for macular edema, have an eye infection, or are being treated for glaucoma with eye drops, unless cleared by your treating ophthalmologist. LATISSE™ is not approved for people under the age of 18. It is not recommended for pregnant or lactating women. You should discontinue use of Latisse and call your physician immediately if you develop an eye infection, sudden decrease in vision, suffer eye trauma, or develop eye or eyelid reactions.

I have read the above information and have discussed the risks, benefits, side effects and alternatives to using Latisse. I understand that it is impossible for the physician to inform me of every possible complication that may occur. My physician has told me that results cannot be guaranteed. I understand that Pelle Spa will not accept returns or issue refunds, unless the product is damaged or defective. By signing below, I agree that my physician has answered all of my questions and I give informed consent to proceed with Latisse treatment.

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Patient Signature Date