

LICALTIL/CVC LICALTILLUCTODY

Dr. Maria Tetelbaum

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LATISSE® Eye Evaluation & Consent Form

HEALIN/ETE HEALIN HISTORY		Date:
First Name (& preferred):	Last Name:	
Date of Birth: D M YR	Age:	Pronouns: She/Her He/Him Other:
Address:	City/Town:	Postal Code:
Telephone: ()	May we lea	ve a voicemail? YES / NO
Email:		Occupation:
OHIP (health card) #:(*Will only be used if medical assessment/eye exam a	nd/or treatment is provid	Version Code:
How did you hear about us?		
Date of last eye exam:	Eye Doctor (name & pho	ne #):
At your last eye examination, were you told that you h	nad any problems with yo	our eyes? Y/N If yes, please explain:
Your eye colour:		
Do you have a history of glaucoma or increased pre	ssure inside the eye?	/ / N
Does anyone in your immediate family have a history	of glaucoma or increase	d pressure inside the eye? Y/N
Do you wear eye glasses or contact lenses? Y/N	Have you had	any injuries/surgeries to the eye/eyelids? Y/N
Do you now, or have you ever, had any visual problem	ns with one or both eyes	? Y/N
Are you bothered by "dry eyes"? Y/N	Do you have overly "wa	tery eyes"? Y/N
Do you have "sensitive eyes"? (i.e.: become irritated e	easily, itchy, red, swell ex	ccessively) Y/N If yes, please describe:
Do you use any eye drops/eye medications of any kin	nd? Y/N If yes,	please explain:
Do you have any allergies /sensitivities? Y/N	If yes, please explain	·
Current health conditions /diagnoses (overall health):		
Please list any current medications including vitamin	ns, herbal supplements (dosage not required)
Are you planning a pregnancy within the next year?	Y / N	
Have you ever used Latisse® or any products markete	ed for lash growth/thicke	ening? Y/N If so, which product?

MEDICATION INFORMATION (*please read)

Latisse® solution (bimatoprost ophthalmic solution) 0.03% is indicated to treat hypotrichosis (inadequate or deficiency of hair) of the eyelashes by increasing their growth, including length, thickness, and darkness. 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 medication is applied once each night along the upper lash line only. The effect of this medication is gradual, but most patients see improvement by 2 months. The effect of this medication is not permanent. If you stop using this medication, your eyelashes will gradually return to their original appearance. Continued use is required to maintain desired results. Results are not guaranteed.

Common side effects after using Latisse® are an itching sensation in the eyes and/or eye redness (reported in approximately 4% of patients). Less common side effects typically occur on the skin close to where Latisse® is applied, or in the eyes. These include eye

irritation, dryness of the eyes, redness of the eyelids, and skin darkening which may be reversible. Latisse® use may also cause increased brown pigmentation of the coloured part of the eye (the iris) which is likely to be permanent. It is possible for hair growth to occur in other areas of your skin that Latisse® frequently touches. There may be other possible side effects/complications not known at this time.

If you develop a new ocular condition (e.g., trauma or infection), experience a sudden decrease in vision, have eye surgery, or develop any reactions, particularly conjunctivitis and eyelid reactions, you should immediately seek your physician's advice concerning the continued use of Latisse®.

You should inform your physician if you are using LATISSE® especially if you have a history of eye pressure problems. If any eye surgery is planned, inform your ophthalmologist that you are using Latisse®. Inform anyone conducting an eye pressure examination that you are using Latisse®.

Do not use Latisse® if you are allergic or hypersensitive to bimatoprost (LumiganR) or any other ingredient in this product.

Latisse® is **not** to be administered if you are pregnant or breastfeeding.

This product is non-returnable.

Please initial (do not check) that you understand and agree with the statements below:			
I do not have glaucoma and I have disclosed my general/eye specific health history to the best of my knowledge I am not pregnant or breastfeeding.			
I am aware Latisse® is a prescription medication and I am responsible for reading the package insert before use. I have been given a copy of the <i>Latisse® Instructions For Use</i> sheet and understand the steps of application.			
I have read and understood ALL of the information provide	d within this 2 page document, and am aware of and accept the		
the physician or nurse.	had sufficient opportunity to discuss my concerns/questions with		
This consent is considered valid for subsequent treatments	s unless revoked in writing.		
Dated:	Dated:		
<u> </u>	Butcu.		
PATIENT Name (printed)	Witness/Provider Name (printed)		
Signature (patient or legal guardian)	Signature		
(patient of legal guardian)			



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Latisse® Instructions For Use

(for patient to keep)

STEP 1. Before Applying

Before applying Latisse® (bimatoprost ophthalmic solution) 0.03% each night, ensure your face is clean, makeup and contact lenses are removed, and any other facial care products have already been applied.

STEP 2. Prepare the Applicator

Remove an applicator from its tray. Then, holding the sterile applicator horizontally, place one drop of Latisse® solution on the area of the applicator closest to the tip but not on the tip.

STEP 3. Apply to Skin Using Applicator

Immediately draw the applicator carefully along the skin at the base of the upper eyelashes (where the eyelashes meet the skin) going from the inner part of your lash line to the outer part. DO NOT APPLY in your eye or to the lower lid.

STEP 4. Blot Excess

The upper lid margin in the area of lash growth should feel lightly moist without runoff. Blot any excess solution runoff outside the upper eyelid margin with a tissue or other absorbent cloth.

STEP 5. Dispose of Applicator

Dispose of the applicator after one use. Repeat for opposite eyelid, using a new sterile applicator to help minimize any potential for contamination from one eyelid to another.

IMPORTANT REMINDERS

- Only use the sterile applicators supplied with Latisse® to apply the product.
- · Apply Latisse® daily for 16 weeks. You should not reduce or stop application when you first notice results.
- Do **not** allow the tip of the bottle or applicator to touch fingers or any other unintended surface, as contamination by common bacteria is known to cause infections.
- · Remove contact lenses prior to applying Latisse®. Contact lenses may be reinserted 15 minutes afterwards.
- Do **not** apply in the eye or to the lower eyelash line.
- It is possible for hair growth to occur on other areas of the skin that Latisse® solution frequently touches, so blot the excess as
 directed.
- · Additional applications of Latisse® will not increase the growth of eyelashes, so please only 1 application per day (at night).
- If you miss a dose, do not play"catch-up." Just apply Latisse® solution the next night as usual.
- · Upon discontinuation of treatment, eyelash growth is expected to return to its pre-treatment level.

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Latisse® is a prescription medication - please read the package insert before use.

*Please contact the clinic immediately if you experience side effects beyond one week or if you develop a new ocular condition (e.g., trauma or infection), experience a sudden decrease in vision, have eye surgery, or develop any reactions, particularly conjunctivitis and eyelid reactions.

Please call 519-266-3642 should you have any concerns.