LATISSE® is a prescription treatment for hypotrichosis of the eyelashes. It is possible for hair growth to occur in other areas of your skin that are not intended. If this occurs, you should stop using LATISSE® and contact your physician.

LATISSE® solution is intended for use on the skin of the upper eyelid margins at the base of the eyelashes. Refer to Illustration 2. DO NOT APPLY to the lower eyelid. If you are using LUMIGAN® or other products in the same class for elevated intraocular pressure (IOP), or if you have a history of abnormal IOP, you should only use LATISSE® under the close supervision of your physician.

LATISSE® use may cause darkening of the eyelid skin which may be reversible. LATISSE® use may also cause increased brown pigmentation of the colored part of the eye which is likely to be permanent.

It is possible for hair growth to occur in other areas of your skin that LATISSE® frequently touches. Any excess solution outside the upper eyelid margin should be wicked up with a tissue or other absorbent material to reduce the chance of this happening. It is also possible for a difference in eyelash length, thickness, fullness, pigmentation, number of eyelash hairs, and/or direction of eyelash growth to occur between eyes. These differences, should they occur, will usually go away if you stop using LATISSE®.

Who should I tell that I am using LATISSE®? You should tell your physician you are using LATISSE® especially if you have a history of eye pressure problems. You should also tell anyone conducting an eye pressure screening that you are using LATISSE®.

What should I do if I get LATISSE® in my eye? LATISSE® solution is an ophthalmic drug product. LATISSE® is not expected to cause harm if it gets into the eye. Do not attempt to rinse your eye in this situation.

What are the possible side effects of LATISSE®? The most common side effect after using LATISSE® solution is an itching sensation in the eyes and/or eye redness. This was reported in approximately 4% of patients. LATISSE® solution may cause other less common side effects which typically occur on the skin close to where LATISSE® is applied, or in the eyes. These include skin darkening, eye irritation, dryness of the eyes, and redness of the eyelids.

If you develop a new ocular condition (e.g., trauma or infection), you should immediately seek your physician's advice. LATISSE® may also cause increased brown pigmentation of the colored part of the eye which is likely to be permanent.

If you miss a dose, don’t try to “catch up.” Just apply LATISSE® solution the next evening. Fifty percent of patients treated with LATISSE® in a clinical study saw significant improvement by 2 months after starting treatment.

Use of LATISSE® more than once a day will not increase the growth of eyelashes more than use once a day. Store LATISSE® solution at 36°F to 77°F (2°C to 25°C).

General Information about LATISSE® Prescription treatments are sometimes prescribed for conditions that are not mentioned in patient information leaflets. Do not use LATISSE® solution for a condition for which it was not prescribed. Do not give LATISSE® to other people. It may not be appropriate for them to use.

This leaflet summarizes the most important information about LATISSE® solution. If you would like more information, talk with your physician. You can also call Allergan’s product information department at 1-800-433-8871.

What are the ingredients in LATISSE®? Active ingredients: bimatoprost Inactive ingredients: borax; disodium edetate; disodium phosphate; sodium chloride; sodium metabisulfite; sodium phoshate; dibasic citric acid; and purified water. Sodium hydroxide and/or hydrochloric acid may be added to adjust pH. The pH during its shelf life ranges from 6.8 – 7.8.

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