PART III: CONSUMER INFORMATION

Sandoz® Latanoprost
Latanoprost Ophthalmic Solution

This leaflet is designed specifically for Consumers. This leaflet is a summary and will not tell you everything about Sandoz Latanoprost. Contact your doctor or pharmacist if you have any questions about the drug.

ABOUT THIS MEDICATION

What the medication is used for:
Sandoz Latanoprost is used to treat ocular hypertension (high pressure in the eye) in patients with open-angle glaucoma or ocular hypertension. These conditions may eventually affect your eyesight.

What it does:
Sandoz Latanoprost is a solution for use only in the eyes. The active ingredient in Sandoz Latanoprost is one of a group of medications known as prostaglandins. It helps to lower the pressure within the eye by increasing the natural outflow of fluid from inside the eye.

When it should not be used:
Do not use Sandoz Latanoprost if you have a known hypersensitivity to benzalkonium chloride or any other ingredient in this product (see "What the medicinal ingredient is." and "What the nonmedicinal ingredients are.").

What the medicinal ingredient is:
Latanoprost.

What the nonmedicinal ingredients are:
Benzalkonium chloride, sodium chloride, sodium dihydrogen phosphate dihydrate, disodium phosphate anhydrous, water for injection, hydrochloric acid and/or sodium hydroxide to adjust pH.

What dosage forms it comes in:
Each bottle contains 2.5 mL of solution, approximately 80 drops. Each millilitre (mL) contains 50 micrograms of latanoprost.

WARNINGS AND PRECAUTIONS

BEFORE you use Sandoz Latanoprost talk to your doctor or pharmacist if:
– You are allergic to any of the ingredients in Sandoz Latanoprost
– You are using any other eye drops or taking any other medication
– You are pregnant, think you might be pregnant or you are planning a pregnancy
– You are breast feeding or planning to breast feed
– You have or have had herpes simplex keratitis (inflammation caused by the herpes simplex virus)
– You have liver or kidney problems
– You have or have had eye inflammation (e.g., uveitis, iritis)

Sandoz Latanoprost contains a preservative that may be absorbed by contact lenses and stains them a brown colour. Contact lenses can be reinserted 15 minutes after applying the eye drops.

If you are using more than one type of eye drop medication, wait at least 5 minutes between each different eye drop.

INTERACTIONS WITH THIS MEDICATION

Studies have shown that precipitation occurs when eye drops containing thimerosal are mixed with Sandoz Latanoprost. If such drugs are used, they should be administered with an interval of at least 5 minutes between applications.

Only on your doctor’s advice, Sandoz Latanoprost may be used concomitantly with other topical ophthalmic products to further lower intraocular pressure. If more than one topical ophthalmic drug is being used, the drugs should be administered at least 5 minutes apart.

PROPER USE OF THIS MEDICATION

Usual adult dose:
One drop of Sandoz Latanoprost should be dropped into the affected eye(s) once daily. The best time to do this is in the evening.

Do not allow the dropper tip of the bottle to touch the eye or other surrounding structures, because this could contaminate the tip with common bacteria known to cause eye infections. Serious damage to the eye with subsequent loss of vision may result if you use eye drop solutions that have become contaminated. If you experience any type of eye condition or have surgery, immediately seek your doctor’s advice concerning the continued use of the bottle you are using.

If you forget to use your eye drops at the usual time, wait until it is time for your next dose. If you put too many drops in your eye(s), you may feel some slight irritation.

Sandoz Latanoprost is not recommended for use in children.

Overdose:
In case of drug overdose, contact a healthcare practitioner, hospital emergency department or regional Poison Control Centre immediately, even if there are no symptoms.

Missed dose:
If one dose is missed, treatment should continue with the next dose the following day.

SIDE EFFECTS AND WHAT TO DO ABOUT THEM

Sandoz Latanoprost may change the colour of your eye. It may make your iris (the coloured part of your eye) more brown. This happens most commonly if your iris has mixed colours, i.e., blue-brown, grey-brown, green-brown or yellow-brown. If you use Sandoz Latanoprost in one eye only, colour changes in the iris may appear only in the treated eye. These changes may be permanent.

Sandoz Latanoprost may also cause your eyelashes to darken, appear thicker and longer than they usually do and increase in number. Sandoz Latanoprost might cause eye irritation due to the growth of misdirected eyelashes; tell your doctor if this happens. A very small number of people may notice their eyelids look darker after using Sandoz Latanoprost for some time. These changes may be more noticeable if you are only treating one eye. Eyelash changes are reversible after treatment with Sandoz Latanoprost is stopped. Eyelid skin darkening may be permanent.

When using Sandoz Latanoprost, you might feel as if there is something in your eye(s). Your eye(s) might water and become red. As with other eye drops, if your vision is blurred when you first put your drops in, wait until this wears off before you drive or operate machinery. A few people using Sandoz Latanoprost have developed a skin rash.
A few people may experience changes in their vision, sometimes in combination with a red and sore/painful eye. These changes do not always occur right after administering the drops, and if they occur, you may find that reading and seeing fine details is more difficult. Although unlikely, if you experience any of these changes, stop using Sandoz Latanoprost and contact your doctor immediately.

Sandoz Latanoprost may cause the following side effects as well.

*Common Ocular Side Effects:* burning and stinging, blurred vision, red eyes, foreign body sensation, itching, increased iris pigmentation, damage of the cornea in a pinpoint pattern, dry eyes, excessive tearing, eye pain, eyelid crusting, red and swollen eyelid, eyelid discomfort/pain, photophobia (visual sensitivity to light).

*Uncommon Ocular Side Effects:* discharge from the eye, diplopia (doubled vision), conjunctivitis, iritis/uveitis (inflammation of the interior of the eye), darkening of the palpebral skin (skin related to the eyelid).

*Common Systemic Side Effects:* upper respiratory tract infection/cold/flu, pain in muscle/joint/back, chest pain/angina pectoris, rash/allergic skin reaction.

Be sure to tell your doctor (or pharmacist) if you notice any other unwanted side effects.

### SERIOUS SIDE EFFECTS, HOW OFTEN THEY HAPPEN AND WHAT TO DO ABOUT THEM

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<th>Talk with your doctor or pharmacist</th>
<th>Stop taking drug and call your doctor or pharmacist</th>
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<td>In all cases</td>
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<td><strong>Uncommon ocula</strong></td>
<td><strong>Uncommon ocula</strong></td>
<td><strong>Systemic Adverse Events</strong></td>
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<td>Macular edema: blurred or wavy vision in the middle of the eye and colour perception changes</td>
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<td>Herpetic keratitis: infection and infestations of the eyes (blurred vision, pain, redness, tearing, discharge, sensitivity to lights)</td>
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<td>Asthma/Asthma aggravation/Acute asthma attack/Difficulty to breathe</td>
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<td>Severe skin reactions including rash and skin degradation in different parts of the body</td>
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This is not a complete list of side effects. For any unexpected effects while taking Sandoz Latanoprost, contact your doctor or pharmacist.

### HOW TO STORE IT

Before Sandoz Latanoprost is first opened, keep it in a fridge (between 2°C and 8°C), out of direct light. Store in the original carton to protect from light. Once the bottle has been opened, Sandoz Latanoprost may be kept at room temperature (between 15°C and 25°C). Sandoz Latanoprost must be used within 28 days after opening the bottle. Discard the bottle and/or unused contents after 28 days. Sandoz Latanoprost should not be used after the expiry date on the bottle.

Keep all medicines in a safe place, out of the reach of children.

### REPORTING SUSPECTED SIDE EFFECTS

You can report any suspected adverse reactions associated with the use of health products to the Canada Vigilance Program by one of the following 3 ways:

- Report online at [www.healthcanada.gc.ca/medeffect](http://www.healthcanada.gc.ca/medeffect)
- Call toll-free at 1-866-234-2345
- Complete a Canada Vigilance Reporting Form and:
  - Fax toll-free to 1-866-678-6789, or
  - Mail to: Canada Vigilance Program
    Health Canada
    Postal Locator 0701E
    Ottawa, Ontario
    K1A 0K9

Postage paid labels, Canada Vigilance Reporting Form and the adverse reaction reporting guidelines are available on the MedEffect™ Canada Web site at [www.healthcanada.gc.ca/medeffect](http://www.healthcanada.gc.ca/medeffect)

**NOTE:** Should you require information related to the management of side effects, contact your health professional. The Canada Vigilance Program does not provide medical advice.

### MORE INFORMATION

This document plus the full product monograph, prepared for health professionals can be found by contacting the sponsor, Sandoz Canada Inc., at:

1-800-361-3062

or by written request at:

145 Jules-Léger

Boucherville, Québec

J4B 7K8 Canada

Or by e-mail at: medinfo@sandoz.com

This leaflet was prepared by Sandoz Canada Inc.

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