

## **INFORMATION**

### **ISOPTO\* CARPINE (pilocarpine hydrochloride) Eye Drops, 1%, 2%, 4%**

## **DESCRIPTION**

Pilocarpine hydrochloride 1%, 2%, and 4%, hypromellose 0.5%, benzalkonium chloride 0.01%, boric acid, sodium citrate and purified water.

## **PHARMACOLOGY**

Lowers intraocular pressure.

## **INDICATIONS**

In the treatment of glaucoma.

## **CONTRAINDICATIONS**

Miotics are contraindicated in conditions where pupillary constriction is undesirable such as in acute iritis or anterior uveitis

Hypersensitivity to the active substance or to any of the excipients.

## **PRECAUTIONS**

Sensitivity is infrequently observed. If reaction occurs, discontinue use.

Retinal detachment has been reported when miotics are used in susceptible individuals, such as young patients with myopia or patients with history of retinal detachment.

Miotics should be avoided in acute inflammatory diseases of the anterior chamber.

A paradoxical rise in intraocular pressure may be observed in patients with severely compromised trabecular outflow.

Caution is advised in the presence of corneal or conjunctival damage to avoid excessive penetration which can produce systemic toxicity.

ISOPTO CARPINE should be used with caution in patients with acute cardiac failure, bronchial asthma, peptic ulcer, hyperthyroidism, gastro-intestinal spasms, Parkinson's Disease, urinary tract obstruction, recent myocardial infarction, hypertension and hypotension due to the risk of exacerbating these conditions.

ISOPTO CARPINE contains benzalkonium chloride which may cause eye irritation and is known to discolour soft contact lenses. Avoid contact with soft contact lenses. Patients must be instructed to remove contact lenses prior to the application of ISOPTO CARPINE and wait at least 15 minutes before reinsertion.

Nasolacrimal occlusion or gently closing the eyelid after administration is recommended. This may reduce the systemic absorption of medicinal products administered via the ocular route and result in a decrease in systemic adverse reactions.

## **Effects on fertility**

Studies have not been performed to evaluate the effect of topical ocular administration of ISOPTO CARPINE on fertility.

## **Use in Pregnancy**

**Category B3** - There are no or limited amount of data from the use of ISOPTO CARPINE in pregnant women. Animal studies have, however, showed harmful effects of systemic pilocarpine exposure with respect to reproductive toxicity in rats. There are no adequate and well controlled studies in pregnant women. As a precautionary measure, it is preferable to avoid the use of ISOPTO CARPINE during pregnancy.

## **Use in Lactation**

It is unknown whether pilocarpine is excreted in human milk. However, excretion in breast milk should be expected. There is also no information on the safety of pilocarpine ophthalmic formulations used during breast feeding. However, a risk to the suckling child cannot be excluded. A decision must be made whether to discontinue breast-feeding or to abstain from ISOPTO CARPINE, taking into account the benefit of breast feeding for the child and the benefit of therapy for the woman.

## **Effects on Ability to Drive and Use Machines**

ISOPTO CARPINE has a major influence on the ability to drive and use machines. Miosis may cause blurred vision and difficulty in dark adaptation. Patients should be advised to exercise caution while driving at night or while performing hazardous tasks in poor light.

## **INTERACTION WITH OTHER MEDICINES**

No interaction studies have been performed.

## **ADVERSE EFFECTS**

The following adverse reactions are classified according to the following convention: very common ( $\geq 1/10$ ), common ( $\geq 1/100$  to  $<1/10$ ), uncommon ( $\geq 1/1,000$  to  $<1/100$ ), rare ( $\geq 1/10,000$  to  $<1/1,000$ ), very rare ( $<1/10,000$ ), or not known (cannot be estimated from the available data) according to system organ classes. Within each frequency-grouping, adverse reactions are presented in order of decreasing seriousness. The adverse reactions have been observed during clinical trials and post-marketing experience with ISOPTO CARPINE.

### ***Nervous system disorders***

Very Common ( $\geq 10\%$ ): headache

Common ( $\geq 1\%$  to  $< 10\%$ ): dizziness.

### ***Eye disorders***

Very Common ( $\geq 10\%$ ): vision blurred

Common ( $\geq 1\%$  to  $< 10\%$ ): visual impairment, visual acuity reduced, eye pain, photopsia, vitreous floaters, myodesopsia, eye irritation, ocular hyperaemia

Uncommon ( $\geq 0.1\%$  to  $< 1\%$ ): retinal tear, vitreous haemorrhage, eyelid oedema, miosis, vitreous detachment, glare, foreign body sensation in eyes

Not Known: intraocular pressure increased, corneal oedema.

### ***Gastrointestinal disorders***

Common ( $\geq 1\%$  to  $< 10\%$ ): nausea

Not Known: vomiting

Slight ciliary spasms may occur with resultant temporary reduction of visual acuity.

## **DOSAGE AND ADMINISTRATION**

Topically, 1-2 drops in the eye(s) 3-4 times daily.

## **OVERDOSAGE**

An ocular overdose of ISOPTO CARPINE can be flushed from the eye(s) with lukewarm water. In case of overdose, symptoms of toxicity may include: headache, salivation, sweating, syncope, bradycardia, hypotension, abdominal cramps, vomiting, asthma and diarrhoea. Treatment of overdose is supportive. In cases of severe systemic toxicity therapy with anticholinergics may be necessary.

Contact the Poisons Information Centre on 13 11 26 for advice on management.

## **PRESENTATION AND STORAGE CONDITIONS**

DROP-TAINER\* 1%, 2%, 4%: 15 mL.

*Consumer Product Information supplied with this product.*

Store below 25°C. Do not freeze. Protect from light.

Discard container 4 weeks after opening.

## **NAME AND ADDRESS OF SPONSOR**

Novartis Pharmaceuticals Australia Pty Limited

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## **POISON SCHEDULE OF THE MEDICINE**

Prescription Only Medicine (Schedule 4)

## **DATE OF FIRST INCLUSION IN THE ARTG**

15 October 1991

## **DATE OF MOST RECENT AMENDMENT**

4 July 2017

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