

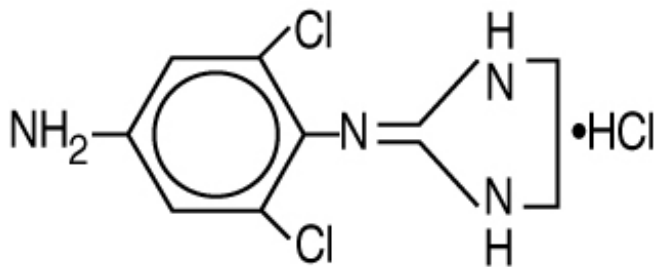
IOPIDINE - apraclonidine hydrochloride solution/ drops
Alcon Laboratories, Inc.

IOPIDINE* 1%
(apraclonidine hydrochloride ophthalmic solution) 1% as base
Sterile

DESCRIPTION

IOPIDINE 1% Ophthalmic Solution contains apraclonidine hydrochloride, an alpha adrenergic agonist, in a sterile isotonic solution for topical application to the eye. Apraclonidine hydrochloride is a white to off-white powder and is highly soluble in water. Its chemical name is 2-[(4-amino-2,6 dichlorophenyl)imino] imidazolidine monohydrochloride with an empirical formula of C₉H₁₁Cl₃N₄ and a molecular weight of 281.6.

The chemical structure of apraclonidine hydrochloride is:



Each mL of IOPIDINE 1% Ophthalmic Solution contains: **Actives:** apraclonidine hydrochloride 11.5 mg equivalent to apraclonidine base 10 mg. **Inactives:** sodium chloride, sodium acetate, sodium hydroxide and/or hydrochloric acid (pH 4.4-7.8), purified water and benzalkonium chloride 0.01% (preservative). Osmolality is 260-320 mOsm.

CLINICAL PHARMACOLOGY

Apraclonidine is a relatively selective, alpha adrenergic agonist and does not have significant membrane stabilizing (local anesthetic) activity. When instilled into the eye, IOPIDINE 1% (apraclonidine hydrochloride ophthalmic solution) has the action of reducing intraocular pressure (IOP). Ophthalmic apraclonidine has minimal effect on cardiovascular parameters.

Optic nerve head damage and visual field loss may result from an acute elevation in IOP that can occur after argon or Nd:YAG laser surgical procedures. Elevated IOP, whether acute or chronic in duration, is a major risk factor in the pathogenesis of visual field loss. The higher the peak or spike of IOP, the greater the likelihood of visual field loss and optic nerve damage especially in patients with previously compromised optic nerves. The onset of action with IOPIDINE 1% Ophthalmic Solution can usually be noted within one hour and the maximum IOP reduction usually occurs three to five hours after application

of a single dose. The precise mechanism of the ocular hypotensive action of IOPIDINE 1% Ophthalmic Solution is not completely established at this time. Aqueous fluorophotometry studies in man suggest that its predominant action may be related to a reduction of aqueous formation. Controlled clinical studies of patients requiring argon laser trabeculoplasty, argon laser iridotomy or Nd:YAG posterior capsulotomy showed that IOPIDINE 1% Ophthalmic Solution controlled or prevented the post-surgical IOP rise typically observed in patients after undergoing those procedures. After surgery, the mean IOP was 1.2 to 4 mmHg below the corresponding pre-surgical baseline pressure before IOPIDINE Ophthalmic Solution treatment. With placebo treatment, post-surgical pressures were 2.5 to 8.4 mmHg higher than their corresponding pre-surgical baselines. Overall, only 2% of patients treated with IOPIDINE* 1% Ophthalmic Solution had severe IOP elevations (spike greater than or equal to 10 mmHg) during the first three hours after laser surgery, whereas 23% of placebo-treated patients responded with severe pressure spikes (Table 1). Of the patients that experienced a pressure spike after surgery, the peak IOP was above 30 mmHg in most patients (Table 2) and was above 50 mmHg in seven placebo-treated patients and one IOPIDINE 1% Ophthalmic Solution-treated patient.

Table 1: Incidence of IOP Spikes Greater Than or Equal to 10 mmHg

Study	Laser Procedure	P-Value	Treatment			
			Apraclonidine		Placebo	
			^a N	(%)	^a N	(%)
1	Trabeculoplasty	<0.05	0/40	(0%)	6/35	(17%)
2	Trabeculoplasty	=0.06	2/41	(5%)	8/42	(19%)
1	Iridotomy	<0.05	0/11	(0%)	4/10	(40%)
2	Iridotomy	=0.05	0/17	(0%)	4/19	(21%)
1	Nd:YAG Capsulotomy	<0.05	3/80	(4%)	19/83	(23%)
2	Nd:YAG Capsulotomy	<0.05	0/83	(0%)	22/81	(27%)

^aN = Number Spikes/Number Eyes.

Table 2: Magnitude of Post-surgical IOP in Trabeculoplasty, Iridotomy and Nd:YAG Capsulotomy Patients With Severe Pressure Spikes Greater than or Equal to 10 mmHg

Maximum Postsurgical IOP (mmHg)

Treatment	Spikes	Total			
		20-29 mmHg	30-39 mmHg	40-49 mmHg	> 50 mmHg
IOPIDINE	8	1	4	2	1
Placebo	78	16	47	8	7

INDICATIONS AND USAGE

IOPIDINE 1% Ophthalmic Solution is indicated to control or prevent post-surgical elevations in IOP that occur in patients after argon laser trabeculoplasty, argon laser iridotomy or Nd:YAG posterior capsulotomy.

CONTRAINDICATIONS

IOPIDINE 1% Ophthalmic Solution is contraindicated for patients receiving monoamine oxidase inhibitor therapy and for patients with hypersensitivity to any component of this medication or to clonidine.

WARNINGS

FOR TOPICAL OPHTHALMIC USE ONLY. Not for injection or oral ingestion.

PRECAUTIONS

General

Since IOPIDINE* 1% Ophthalmic Solution is a potent depressor of IOP, patients who develop exaggerated reductions in IOP should be closely monitored. Although the acute administration of two drops of IOPIDINE 1% Ophthalmic Solution has minimal effect on heart rate or blood pressure in clinical studies evaluating patients undergoing anterior segment laser surgery, the preclinical pharmacologic profile of this drug suggests that caution should be observed in treating patients with severe cardiovascular disease including hypertension. IOPIDINE 1% Ophthalmic Solution should also be used with caution in patients with severe coronary insufficiency, recent myocardial infarction, cerebrovascular disease, chronic renal failure, Raynaud's disease or thromboangiitis obliterans.

The possibility of a vasovagal attack occurring during laser surgery should be considered and caution used in patients with history of such episodes.

Topical ocular administration of two drops of 0.5%, 1%, and 1.5% IOPIDINE Ophthalmic Solution to New Zealand Albino rabbits three times daily for one month resulted in sporadic and transient instances of minimal corneal cloudiness in the 1.5% group only. No histopathological changes were noted in those eyes. No adverse ocular effects were observed in cynomolgus monkeys treated with two drops of 1.5% IOPIDINE Ophthalmic Solution applied three times daily for three months. No corneal changes were observed in 320 humans given at least one dose of IOPIDINE 1% Ophthalmic Solution.

Information for Patients

Apraclonidine can cause dizziness and somnolence. Patients who engage in hazardous activities requiring mental alertness should be warned of the potential for a decrease in mental alertness on the day of surgery.

Drug Interactions

Interactions with other agents have not been investigated.

Carcinogenesis, Mutagenesis, Impairment of Fertility

No significant change in tumor incidence or type was observed following two years of oral administration of apraclonidine HCl to rats and mice at dosages of 1 and 0.6 mg/kg/day, up to 50 and 30 times, respectively, the maximum dose recommended for human topical ocular use. Apraclonidine HCl was not mutagenic in a series of *in vitro* mutagenicity tests, including the Ames test, a mouse lymphoma forward mutation assay, a chromosome aberration assay in cultured Chinese hamster ovary (CHO) cells, a sister chromatid exchange assay in CHO cells, and a cell transformation assay. An *in vivo* mouse micronucleus assay conducted with apraclonidine HCl also provided no evidence of mutagenicity. Reproduction and fertility studies in rats showed no adverse effect on male or female fertility at a dose of 0.5 mg/kg/day (25 times the maximum recommended human dose).

Pregnancy

Apraclonidine HCl has been shown to have an embryocidal effect in rabbits when given in an oral dose of 3 mg/kg/day (150 times the maximum recommended human dose). Dose related maternal toxicity was observed in pregnant rats at 0.3 mg/kg/day (15 times the maximum recommended human dose). There are no adequate and well controlled studies in pregnant women. IOPIDINE* 1% Ophthalmic Solution should be used during pregnancy only if the potential benefit justifies the potential risk to the fetus.

Nursing Mothers

It is not known if topically applied IOPIDINE 1% Ophthalmic Solution is excreted in human milk. Decision should be made to discontinue nursing temporarily for the one day on which IOPIDINE 1% Ophthalmic Solution is used.

Pediatric Use

Safety and effectiveness in pediatric patients have not been established.

Geriatric Use

No overall differences in safety or effectiveness have been observed between elderly and younger patients.

ADVERSE REACTIONS

The following adverse events, occurring in less than 2% of patients, were reported in association with the use of IOPIDINE 1% Ophthalmic Solution in laser surgery: ocular injection, upper lid elevation, irregular heart rate, nasal decongestion, ocular inflammation, conjunctival blanching, and mydriasis.

The following adverse events were observed in investigational studies dosing IOPIDINE 1% Ophthalmic Solution once or twice daily for up to 28 days in non-laser studies:

Ocular

Conjunctival blanching, upper lid elevation, mydriasis, burning, discomfort, foreign body sensation, dryness, itching, hypotony, blurred or dimmed vision, allergic response, conjunctival microhemorrhage.

Gastrointestinal

Abdominal pain, diarrhea, stomach discomfort, emesis

Cardiovascular

Bradycardia, vasovagal attack, palpitations, orthostatic episode

Central Nervous System

Insomnia, dream disturbances, irritability, decreased libido.

Other

Taste abnormalities, dry mouth, nasal burning or dryness, headache, head cold sensation, chest heaviness or burning, clammy or sweaty palms, body heat sensation, shortness of breath, increased pharyngeal secretion, extremity pain or numbness, fatigue, paresthesia, pruritus not associated with rash.

Clinical Practice

The following events have been identified during postmarketing use of IOPIDINE 1% Ophthalmic Solution in clinical practice. Because they are reported voluntarily from a population of unknown size, estimates of frequency cannot be made. The events, which have been chosen for inclusion due to either their seriousness, frequency of reporting, possible causal connection to IOPIDINE 1% Ophthalmic Solution, or a combination of these factors, include hypersensitivity.

OVERDOSAGE

Ingestion of IOPIDINE* 0.5% Ophthalmic Solution has been reported to cause bradycardia, drowsiness, and hypothermia. Accidental or intentional ingestion of oral clonidine has been reported to cause apnea, arrhythmias, asthenia, bradycardia, conduction defects, diminished or absent reflexes, dryness of the mouth, hypotension, hypothermia, hypoventilation, irritability, lethargy, miosis, pallor, respiratory depression, sedation or coma, seizure, somnolence, transient hypertension, and vomiting. Treatment of an oral overdose includes supportive and symptomatic therapy; a patent airway should be maintained. Hemodialysis is of limited value since a maximum of 5% of circulating drug is removed.

DOSAGE AND ADMINISTRATION

One drop of IOPIDINE* 1% Ophthalmic Solution should be instilled in the scheduled operative eye one hour before initiating anterior segment laser surgery and a second drop should be instilled to the same eye immediately upon completion of the laser surgical procedure. Use a separate container for each single-drop dose and discard each container after use.

HOW SUPPLIED

IOPIDINE 1% Ophthalmic Solution as base is a sterile, isotonic, aqueous solution containing apraclonidine hydrochloride.

Supplied as follows: 0.1 mL in plastic ophthalmic dispensers, packaged two per pouch. These dispensers are enclosed in a foil overwrap as an added barrier to evaporation.

0.1 mL (packaged two per pouch) **NDC** 0065-0660-10

Storage: Store at 2°C to 25°C (36°F-77°F).

Protect from light.

Distributed by:

ALCON LABORATORIES, INC.

Fort Worth, Texas 76134 USA

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PRINCIPAL DISPLAY PANEL

NDC 0065-0660-10

IOPIDINE®

(apraclonidine hydrochloride ophthalmic solution) 1% as base

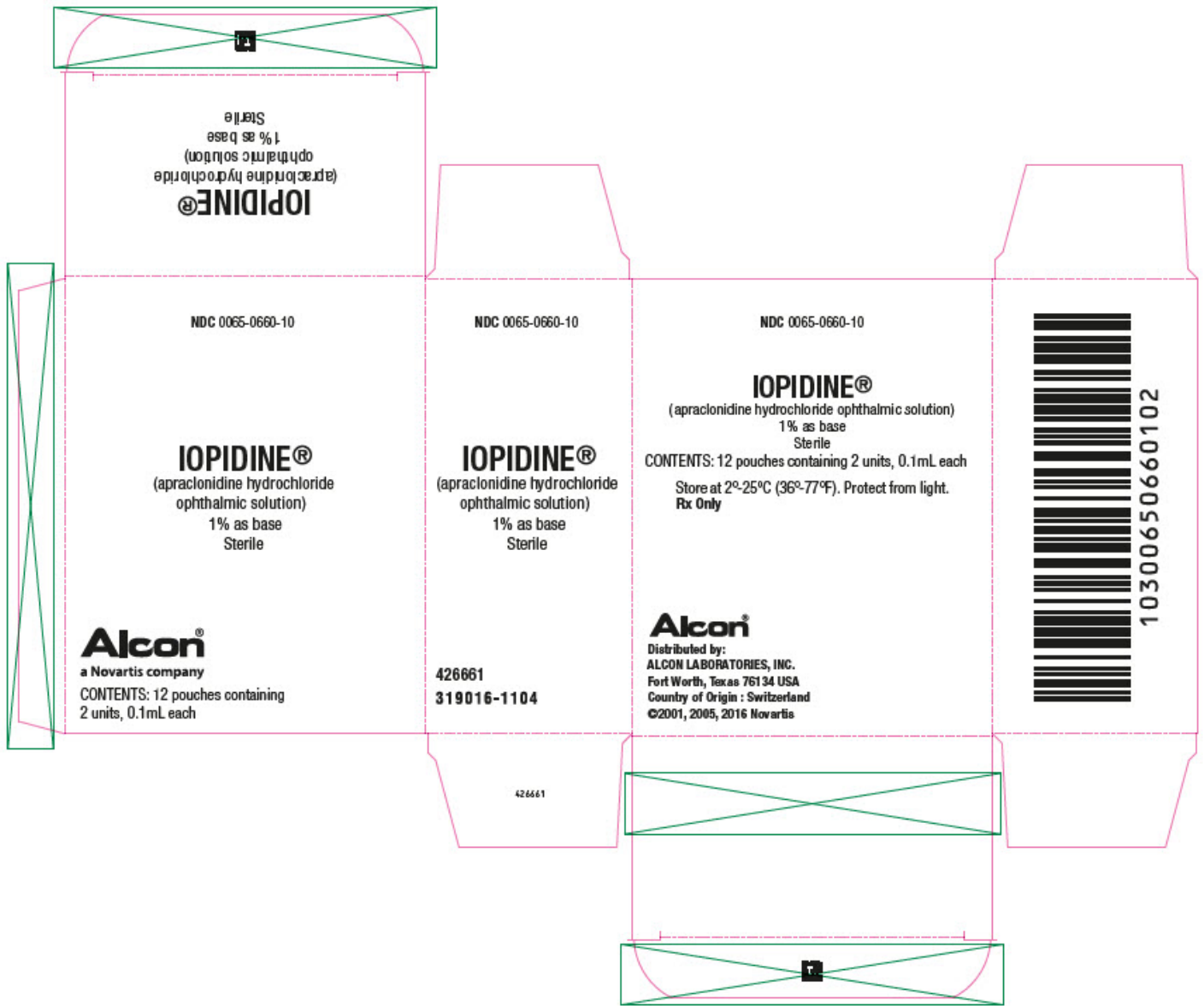
Sterile

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CONTENTS: 12 pouches containing

2 units, 0.1mL each



IOPIDINE

apraclonidine hydrochloride solution/ drops

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:0065-0660
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
APRACLONIDINE HYDROCHLORIDE (UNII: D2VW67N38H) (APRACLONIDINE - UNII:843CEN85DI)	APRACLONIDINE	10 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM ACETATE (UNII: 4550K0SC9B)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0065-0660-10	12 in 1 CARTON	05/15/1988	
1		2 in 1 POUCH		
1		0.1 mL in 1 VIAL, SINGLE-USE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA019779	05/15/1988	

Labeler - Alcon Laboratories, Inc. (008018525)

Revised: 10/2021

Alcon Laboratories, Inc.