

Summary of Product Characteristics

1 NAME OF THE MEDICINAL PRODUCT

Mydrilate 1.0 % w/v Eye Drops

2 QUALITATIVE AND QUANTITATIVE COMPOSITION

Cyclopentolate Hydrochloride 1.0 % w/v.

Contains 10mg cyclopentolate hydrochloride per 1ml

Excipients: also includes benzalkonium chloride solution 0.02% w/v

For a full list excipients, see section 6.1.

3 PHARMACEUTICAL FORM

Eye Drop Solution. (Eye Drops).

Clear colourless or almost colourless liquid.

4 CLINICAL PARTICULARS

4.1 Therapeutic Indications

- (i) Diagnostic purposes for fundoscopy and cycloplegic refraction.
- (ii) Dilating the pupil in inflammatory conditions of the iris and uveal tract.

4.2 Posology and method of administration

Posology

(i) Refraction / Fundoscopy

Adults (and the elderly):

One drop of 0.5% w/v solution instilled into the eye, repeated after 15 minutes if necessary, approximately 40 minutes before examination.

Deeply pigmented eyes may require the use of a 1% w/v solution. NB: maximum effect is reached after 30-60 minutes.

Children 6-16 years:

One drop of 1% w/v solution instilled into the eye, repeated after 15 minutes if necessary, approximately 40 minutes before examination.

Children under 6 years:

One or two drops of 1% w/v solution instilled into the eye, repeated after 15 minutes if necessary, approximately 40 minutes before examination.

(ii) For Uveitis, Iritis and Iridocyclitis

Adults and the elderly:

One or two drops of 0.5% w/v solution instilled into the eye up to 4 times daily or as required.

Deeply pigmented eyes may require the use of a 1% w/v solution.

Children:

At the discretion of the physician.

Paediatric population

Do not use during the first three months of life due to possible association between the cycloplegia produced and the development of amblyopia and also the increased risks of systemic toxicity in neonates.

The lowest possible dose and number of eye drops should be administered in infants. Close monitoring and supervision after instillation is advised.

Elderly population

No specific information on the use of this product in the elderly is available. Clinical trials have included patients over 65 years and no adverse reactions specific to this age group have been reported.

Method of administration

Instill directly into the eye.

To reduce systemic absorption the lacrimal sac should be compressed at the medial canthus by digital pressure for at least two minutes after instillation of the drops. This is especially advisable in children.

4.3 Contraindications

- (i) Use in narrow-angle glaucoma or those with a tendency towards glaucoma e.g. patients with a shallow anterior chamber.
- (ii) Hypersensitivity to cyclopentolate hydrochloride, benzalkonium chloride or any other components of the formulation listed in section 6.1.
- (iii) This preparation contains benzalkonium chloride and should not be used whilst soft contact lenses are being worn.
- (iv) Use in patients with paralytic ileus.
- (v) Use in children with organic brain syndromes, including congenital or neuro- developmental abnormalities, particularly those predisposing to epileptic seizures.

4.4 Special warnings and precautions for use

Because of the risk of precipitating angle-closure glaucoma in the elderly and others prone to raised intraocular pressure, an estimate of the depth of the anterior chamber should be made before use, particularly if therapy is likely to be intense or protracted.

Caution should be observed when drugs of this group are administered to patients with prostatic enlargement, coronary insufficiency or cardiac failure, or ataxia. Atropine-like effects have been reported as side-effects.

Extreme caution is advised for use in children and individuals susceptible to belladonna alkaloids because of the increased risk of systemic toxicity.

Infants appear more likely to absorb eye drops into systemic circulation due to lower tear volume capacity. The risk of toxicity is further increased due to low body mass and immature cardiovascular and nervous systems. Use of mydriatic agents has been associated in preterm infants with increased difficulty feeding, abdominal distention, increased gastric aspirate and rare cases of necrotising enterocolitis. Convulsions in children have also been reported (see section 4.8).

Patients should be warned of the oral toxicity of this preparation, and advised to wash their hands after use. If accidentally swallowed, patients should be advised to seek medical attention.

Use with caution in an inflamed eye as the hyperaemia greatly increases the rate of systemic absorption through the conjunctiva.

4.5 Interaction with other medicinal products and other forms of interaction

The effects of anti-muscarinic agents may be enhanced by the concomitant administration of other drugs with antimuscarinic properties such as some antihistamines, butyrophenones, phenothiazines, tricyclic antidepressants and amantadine.

4.6 Fertility, pregnancy and lactation

There is insufficient evidence as to drug safety in pregnancy and lactation. This product should not be used during pregnancy unless it is considered essential by a physician.

4.7 Effects on ability to drive and use machines

May cause blurred vision, difficulty in focusing and sensitivity to light. Patients should be warned not to drive or engage in other hazardous activities (including climbing ladders and scaffolding) unless vision is clear. Complete recovery from the effects of Mydrilate Eye Drops may take up to 24 hours.

4.8 Undesirable effects

Local irritation may occur following the use of this product.

Systemic toxicity, though uncommon, may follow administration of a 1% solution. Children are more susceptible than adults. Toxicity is usually transient and may manifest as CNS disturbances. Convulsions have been reported in children, however, based on the available data the frequency could not be determined.

For assessment of adverse reactions (ADRs) the following terms regarding frequency are used:

Very common ($\geq 1/10$); common ($\geq 1/100$ and $< 1/10$); uncommon ($\geq 1/1,000$ and $< 1/100$); rare ($\geq 1/10,000$ and $< 1/1,000$); very rare ($< 1/10,000$); Frequency not known (cannot be estimated from the available data).

Psychiatric Disorders:

Not known: abnormal behaviour, psychotic disorders

Nervous system disorders:

Not known: dizziness, partial seizures, convulsions, loss of consciousness, coma

Eye disorders:

Not known: eye oedema, eye pain, photophobia, mydriasis, eye irritation (stinging), ocular hyperaemia, conjunctivitis

Cardiac disorders:

Not known: bradycardia, tachycardia, palpitations, arrhythmia.

Vascular disorders:

Not known: flushing

Gastrointestinal disorders:

Not known: dry mouth, vomiting, gastrointestinal hypomotility and constipation,

Skin and subcutaneous tissue disorders:

Not known: dry skin

Renal and urinary disorders:

Not known: urinary urgency, urinary retention, dysuria

*General disorders and administration site conditions:**Not known:* gait disturbance*Investigations:**Not known:* increased intraocular pressure*Paediatric population:*

Cases of necrotising enterocolitis and abdominal distention have been reported in infants. However, the frequency of these events is unknown. Cardiopulmonary failure and skin rashes have been reported in the paediatric population. The frequency of the latter events is not known.

Cases of convulsions have also been reported in children. The frequency is not known, though infants and younger children appear to be more at risk.

Reporting of suspected adverse reactions

Reporting suspected adverse reactions after authorisation of the medicinal product is important. It allows continued monitoring of the benefit/risk balance of the medicinal product. Healthcare professionals are asked to report any suspected adverse reactions via HPRA Pharmacovigilance, Earlsfort Terrace, IRL – Dublin 2; Tel + 353 1 6764971; fax +353 1 6762517. Website: www.hpra.ie; E-mail: medsafety@hpra.ie.

4.9 Overdose

Systemic toxicity may occur following topical use, particularly in children. It is manifested by flushing and dryness of the skin (a rash may be present in children), blurred vision, a rapid and irregular pulse, fever, abdominal distension in infants, convulsions and hallucinations and the loss of neuromuscular co-ordination.

Treatment is supportive (there is no evidence that physostigmine is superior to supportive management). In infants and small children the body surface must be kept moist. If accidentally ingested, induce emesis or perform gastric lavage.

5 PHARMACOLOGICAL PROPERTIES**5.1 Pharmacodynamic properties**

Cyclopentolate is an anti-muscarinic agent used topically in the eye as a mydriatic and cycloplegic. The effects are similar to those of atropine, but with a more rapid onset and a shorter duration of action.

5.2 Pharmacokinetic properties

Approximately 80% of any ocularly administered drops (26 microliter) is lost via the nasolacrimal duct, conjunctival blood vessels or spillage. Absorption via the nasolacrimal duct and conjunctival blood vessels occurs directly into the systemic circulation and bypasses initial entero-hepatic metabolism.

Cycloplegia following administration is quick in onset and short-lived. Maximal cycloplegia is achieved within 15-45 minutes of instillation and lasts on average about 20 minutes. Recovery normally takes place in about 4 hours, but very occasionally some effect persists for up to 24 hours.

Mydriasis is produced very rapidly and an average pupil diameter of 7 mm is usually reached 15-30 minutes after instillation of one drop of 0.5% solution. Complete recovery from the mydriatic effect generally occurs spontaneously in not more than 20 hours.

5.3 Preclinical safety data

Nothing stated.

6 PHARMACEUTICAL PARTICULARS

6.1 List of excipients

Boric Acid
Potassium Chloride
Benzalkonium Chloride Solution
Purified Water

6.2 Incompatibilities

Not applicable

6.3 Shelf life

Unopened: 2 years.
Once opened: 28 days.

6.4 Special precautions for storage

Store in a refrigerator (2°C-8°C). Do not freeze. Store in the original package. Do not dilute or dispense from any other container than original bottle.

6.5 Nature and contents of container

5ml dropper bottle of 1.0% w/v solution.
Bottle: LDPE, natural colour
Cap: Melamine, white colour

6.6 Special precautions for disposal of a used medicinal product or waste materials derived from such medicinal product and other handling of the product

When using the product for the first time, screw down the cap firmly on the bottle to pierce the seal at the tip of the plastic nozzle and unscrew the cap for use.

7 MARKETING AUTHORISATION HOLDER

Intrapharm Laboratories Limited
The Courtyard Barns
Choke Lane
Cookham Dean
Maidenhead
Berks, SL6 69T
United Kingdom

8 MARKETING AUTHORISATION NUMBER

PA0997/004/002

9 DATE OF FIRST AUTHORISATION/RENEWAL OF THE AUTHORISATION

Date of first authorisation: 1st April 1979
Date of last renewal: 3rd February 2007

10 DATE OF REVISION OF THE TEXT

May 2015