

HYPROMELLOSE EYE DROPS 0.7%- hypromellose eye drops 0.7% for solution

COVALENT MEDICAL

Active Ingredient

Hypromellose 0.7% BP w/v

DIRECTIONS FOR USE

- Instill 1 or 2 drops in the affected eye, as needed

INACTIVE INGREDIENT

1. Benzalkonium Chloride
2. Borax
3. Boric acid
4. EDTA disodium salt
5. Potassium chloride
6. Purified water
7. Sodium chloride

Tamper Protection

- For your protection a tamper evident ring is attached to the bottle cap
- Upon opening, this will separate from the cap and can be discarded
- Use only if this ring is present and attached when the bottle is first opened

USE

- For use as a lubricant to prevent further irritation or to relieve dryness of the eye

Questions

Call: 1-800-103-7321

Email: info@aurolab.com

Web: www.aurolab.com

KEEP OUT OF REACH OF CHILDREN

- If swallowed, get medical help or contact a poison control center right away

ASK DOCTOR

- If you experience eye pain
- Change in vision
- Continued redness (or) irritation of the eye

- Condition worsens or persists for more than 72 hours

DO NOT USE

- If you are sensitive to any ingredient in this product
- If solution changes color or becomes cloudy

Dosage

Instill 1 or 2 drops in the affected eyes as needed

Warnings

For External Use Only

Indications and Usage

For use as a lubricant to prevent further irritation or to relieve dryness of the eye

Eye Lubricant

Eye Lubricant

CARTON LABEL

Drug Facts

Active Ingredient	Purpose
Hypromellose BP 0.7% w/v..... Lubricant	

Use

- For use as a lubricant to prevent further irritation or to relieve dryness of the eye

Warnings

For external use only

Do not use

- If you are sensitive to any ingredient in this product
- If solution changes color or becomes cloudy

When using this product

- Do not touch the nozzle tip to any surface since this may contaminate the solution
- Remove contact lenses before use
- Replace cap after using

Stop use and ask a doctor if

- You experience eye pain
- Change in vision
- Continued redness (or) irritation of the eye
- Condition worsens or persists for more than 72 hours

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FOCUS PI
Hypromellose Eye Drops BP
0.7% w/v

**POST-INJECTION
EYE DROPS**

LONG LASTING RELIEF
Clinically proven to lubricate, hydrate, & relieve the eyes from irritation.

**10ML
STERILE**
Contains one 10 mL bottle

Drug Facts (continued)

Directions

- Instill 1 or 2 drops in the affected eyes as needed

Other information

- Store below 59°-86°F (15°-30°C)

Inactive Ingredients

Benzalkonium Chloride, Borax, Boric acid, Disodium Edetate, Potassium chloride, Purified water, Sodium chloride

Keep out of reach of children.

If swallowed get medical help or contact a Poison Control Center right away.

For your protection a tamper evident ring is attached to the bottle cap. Upon opening, this ring will separate from the cap and can be discarded. Use only if this ring is present and attached when the bottle is first opened

Questions?

Call: 1-855-663-6287
E-mail: info@focusvitamins.com
Web: www.focusvitamins.com

Manufactured for:
Covalent Management, Inc.
24600 Millstream Drive Suite 410
Aldie, VA 20105
Made in India Code:TN/Drugs/TN00002387

FOCUSVITAMINS.COM



FOCUS PI
Hypromellose Eye Drops BP
0.7% w/v

**POST-INJECTION
EYE DROPS**

LONG LASTING RELIEF
Clinically proven to lubricate, hydrate, & relieve the eyes from irritation.

**10ML
STERILE**
Contains one 10 mL bottle
NDC Code: 82724-007-19

HYPROMELLOSE EYE DROPS 0.7%

hypromellose eye drops 0.7% for solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82724-007
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYPROMELLOSE 2910 (4000 MPA.S) (UNII: RN3152OP35) (HYPROMELLOSE 2910 (4000 MPA.S) - UNII:RN3152OP35)	HYPROMELLOSE 2910 (4000 MPA.S)	7 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
EDETATE DISODIUM (UNII: 7FLD91C86K)	
BORIC ACID (UNII: R57ZHV85D4)	
WATER (UNII: 059QF0KO0R)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82724-007-19	10 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	09/01/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M018	09/01/2022	

Labeler - COVALENT MEDICAL (112467517)

Registrant - AUROLAB (677319965)

Establishment

Name	Address	ID/FEI	Business Operations
AUROLAB		677319965	manufacture(82724-007)

Revised: 1/2024

COVALENT MEDICAL