

**OCTIQ LUBRICANT EYE DROPS- dextran 70 hypromellose solution/ drops
Innovus Pharmaceuticals, Inc.**

Octiq Lubricant Eye Drops

Active ingredients

Dextran 70 0.1%

Hypromellose 2910 0.3%

Purpose

Lubricant

Lubricant

Uses

- for the temporary relief of burning and irritation due to dryness of the eye
- for the temporary relief of discomfort due to minor irritations of the eye or to exposure to wind or sun
- as a protectant against future irritation

Warnings

For external use only

Do not use

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

When using this product

- to avoid contamination, do not touch tip of container to any surface
- replace cap after using

Stop use and ask a doctor if

- you feel eye pain
- changes in vision occur redness or irritation of the eye gets worse or lasts more than 72 hours

Keep out of reach of children

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

Directions

- Put 1 or 2 drops in the affected eye(s) as needed

Other Information

- Store at room temperature

Inactive ingredients

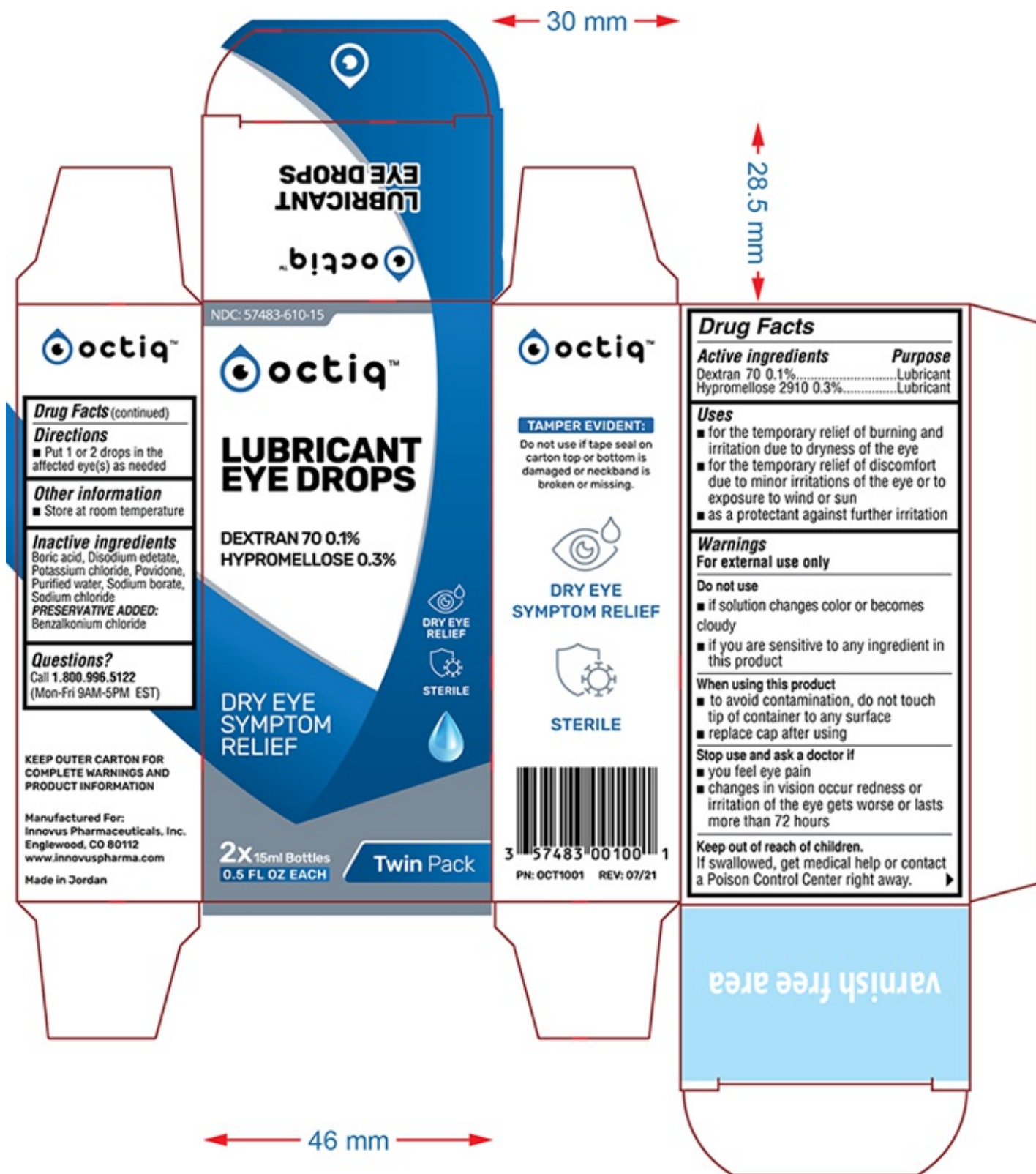
Boric acid, disodium edetate, potassium chloride, povidone, purified water, sodium borate, sodium chloride

Preservative added: benzalkonium choride

Questions?

Call 1.800.996.5122

(Mon-Fri 9AM-5PM EST)



OCTIQ LUBRICANT EYE DROPS

dextran 70 hypromellose solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:57483-610
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTRAN 70 (UNII: 7SA290YK68) (DEXTRAN 70 - UNII:7SA290YK68)	DEXTRAN 70	1 mg in 1 mL
HYPROMELLOSE 2910 (4000 MPA.S) (UNII: RN3152OP35) (HYPROMELLOSE 2910 (4000 MPA.S) - UNII:RN3152OP35)	HYPROMELLOSE 2910 (4000 MPA.S)	3 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BORIC ACID (UNII: R57ZHV85D4)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
POVIDONE (UNII: FZ989GH94E)	
WATER (UNII: 059QF0K00R)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:57483-610-15	2 in 1 CARTON	11/14/2021	
1		15 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M	11/14/2021	

Labeler - Innovus Pharmaceuticals, Inc. (962507187)**Establishment**

Name	Address	ID/FEI	Business Operations
AMMAN PHARMACEUTICAL INDUSTRIES		534677849	manufacture(57483-610)

Revised: 12/2023

Innovus Pharmaceuticals, Inc.