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

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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

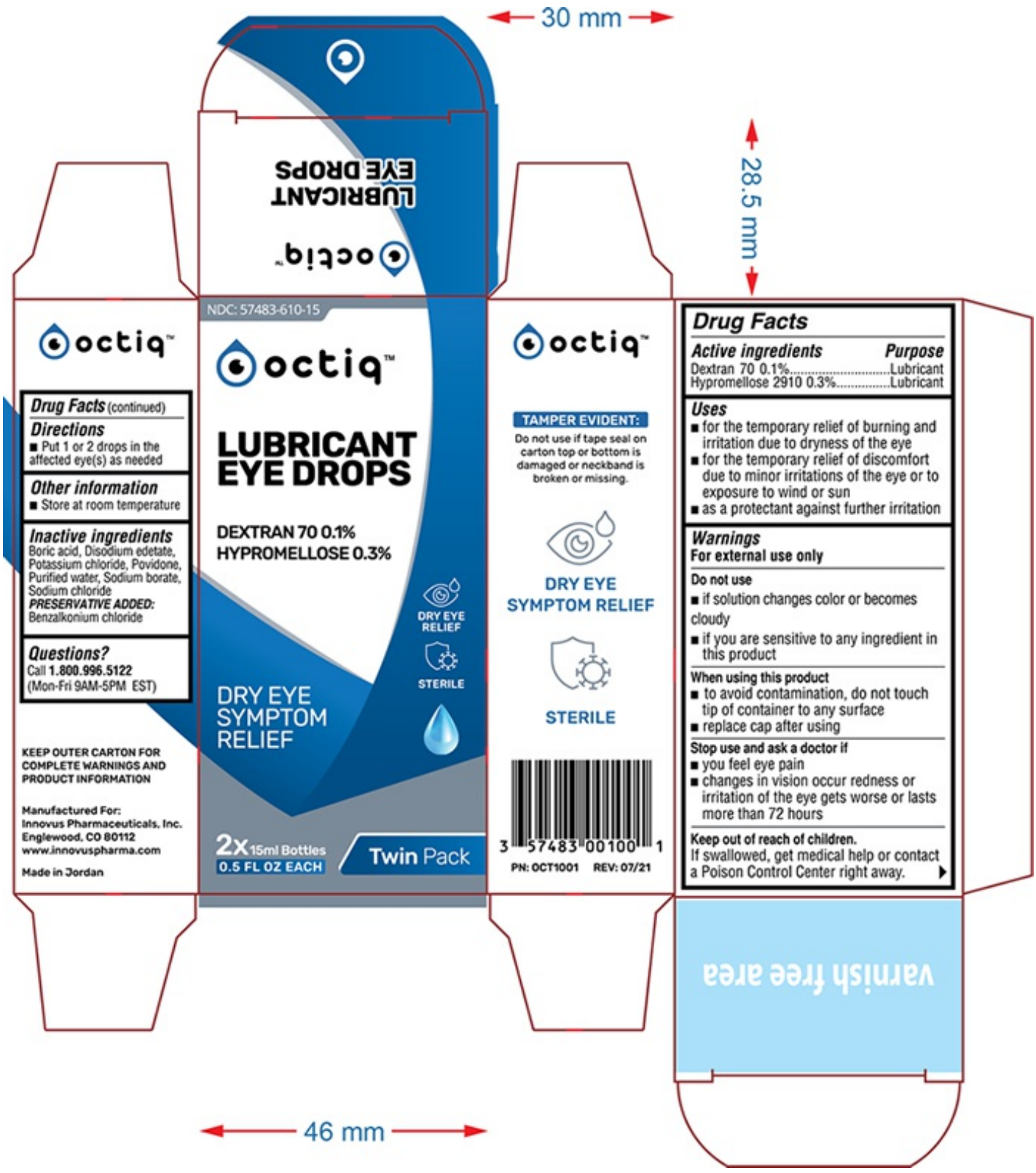
Inactive ingredients

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: 059QF0KO0R)	
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 final	part349	11/14/2021	

Labeler - Innovus Pharmaceuticals, Inc. (962507187)

Establishment

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

Revised: 12/2022 Innovus Pharmaceuticals, Inc.