

MAJOR NATURAL BALANCE- dextran 70, and hypromellose 2910 solution/ drops
Major Pharmaceuticals

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.

Active ingredients	Purpose
Dextran 70 0.1%.....	Lubricant
Hypromellose 2910 0.3%.....	Lubricant

Uses

- relieves dryness of the eye
- prevents further irritation

Warnings

Do not use

- if solution changes color or becomes cloudy

When using this product

- do not touch tip of container to any surface to avoid contamination
- replace cap after each use
- remove contact lenses before using

Stop use and ask a doctor

- if you experience eye pain, changes in vision, continued redness or irritation of the eye
- condition worsens or persists for more than 72 hours

Keep out of reach of children

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

Directions

- instill 1 to 2 drops in the affected eye(s) as needed

Other information

- store at 15°-25°C (59-77°F)
- keep tightly closed

Inactive ingredients

benzalkonium chloride, boric acid, edetate disodium, potassium chloride, purified water, sodium borate, sodium chloride

Distributed by:

Major Pharmaceuticals

31778 Enterprise Drive

Livonia, MI 48150 USA

Made in Korea



MAJOR NATURAL BALANCE

dextran 70, and hypromellose 2910 solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0904-6493
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTRAN 70 (UNII: 7SA290 YK68) (DEXTRAN 70 - UNII:7SA290 YK68)	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
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BORIC ACID (UNII: R57ZHV85D4)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0K00R)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0904-6493-35	1 in 1 CARTON		
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	09/17/2015	

Labeler - Major Pharmaceuticals (191427277)

Revised: 9/2015

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