DROP DREAM EYE 2- polyhexamethylene biguanide liquid K&J.C Co., Ltd

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Drug Facts

polyhexamethylene biguanide
poloxamer407, borax, oric acid, sodium hyaluronate, water, etc
cleanses, rinses, disinfects, stores, lubricates and removed protein buildup
keep out of reach of the children
always use fresh solution to disinfacts, to not reuse solution
keep in room temperature
opthalmic use only



ACTIVE INGREDIENT: Polyhexamecylene Biguanide (20%)

WARNING:

- Always utilize fresh solution for disinfection purposes. Do not reuse the solution.
- This solution is intended for external use only.
 Donot ingest or apply directly to the eyes,
 If you are allergic to any ingredient in this product, DO NOT USE

DIRECTIONS:

- During wear, put 2 or 3 drops on your contacts whenever needed.
- 2. Blink several times. If the contacts still do not feel comfortable, add another drop.
- If discomfort persists, IMMEDIATELY remove the contacts and contact your eye care practitioner.



NDC: 52345-6001-1





Moisturizes Dry Lenses



UP TO 10 HOURS OF MOISTURE



0.44 FL OZ (13mL) STERILE

SAFETY:

Preservative Free

The inner container can be divided into two sections, before and after opening, to ensure solution safety and prevent contamination from external sources.

STORAGE INSTRUCTIONS:

- Keep in a sealed container at room temperature away from direct sublight.
- Discard solution 60 days after opening
- Keep out of reach of children

MANUFACTURER:

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DROP DREAM EYE 2

polyhexamethylene biguanide liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:52345-6001

Route of Administration OPHTHALMIC, NASAL

Active Ingredient/Active Moiety

Ingredient Name

Basis of Strength

Strength

Output

Description: Description of the Strength of the Strength output

Basis of Strength output

Description of the Strength output

Description output

Descr

POLYAMINOPROPYL BIGUANIDE (UNII: DT9D8Z79ET) (POLYAMINOPROPYL BIGUANIDE - UNII:DT9D8Z79ET)

POLYAMINOPROPYL 0.001 mg In 100 mL

Inactive Ingredients

Ingredient Name Strength

WATER (UNII: 059QF0KO0R)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:52345- 6001-1	13 mL in 1 CONTAINER; Type 0: Not a Combination Product	04/27/2024			
Marketing Information						
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
	approved drug her		04/27/2024			

Labeler - K&J.C Co., Ltd (690257639)

Registrant - K&J.C Co., Ltd (690257639)

Establishment						
Name	Address	ID/FEI	Business Operations			
K&J.C Co., Ltd		690257639	manufacture(52345-6001)			

Revised: 5/2024 K&J.C Co., Ltd