**WALGREENS LUBRICANT EYE DROPS PRESERVATIVE FREE- dextran 70, hypromellose liquid**
**Walgreen Company**

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**Walgreens Lubricant Eye Drops Preservative Free (PLD)**

## Active ingredients

Dextran 70 0.1%

Hypromellose 0.3%

## Purposes

Lubricant

Lubricant

## Use

- temporary relief of discomfort due to dryness of the eye or to exposure to wind or sun

## Warnings

For external use only

## Do not use

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

## When using this product

- to avoid contamination, do not touch tip of container to any surface
- do not reuse
- once opened, discard

## Stop use and ask a doctor if

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

## If pregnant or breast-feeding,

ask a health professional before use.

## Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

**Directions**

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

**Other information**

store at 15°-25°C (59°-77°F)

**Inactive ingredients**

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

**Questions or comments?**

Call 1-888-527-4276

**Walgreens Lubricant Eye Drops Preservative Free 35ct**


# WALGREENS LUBRICANT EYE DROPS PRESERVATIVE FREE

dextran 70, hypromellose liquid

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:0363-0646 |
| **Route of Administration** | OPHTHALMIC | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **DEXTRAN 70** (UNII: 7SA290YK68) (DEXTRAN 70 - UNII:7SA290YK68) | DEXTRAN 70 | 0.1 g in 100 mL |
| **HYPROMELLOSE, UNSPECIFIED** (UNII: 3NXW29V3WO) (HYPROMELLOSE, UNSPECIFIED - UNII:3NXW29V3WO) | HYPROMELLOSE, UNSPECIFIED | 0.3 g in 100 mL |

## Inactive Ingredients

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

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:0363-0646-01 | 35 in 1 BOX | 01/01/2020 | |
| **1** | | 0.4 mL in 1 VIAL, SINGLE-USE; Type 0: Not a Combination Product | | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC Monograph Drug | M018 | 01/01/2020 | |

## Labeler - Walgreen Company (008965063)

## Registrant - KC Pharmaceuticals, Inc. (174450460)

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| KC Pharmaceuticals, Inc. | | 174450460 | pack(0363-0646) , label(0363-0646) |

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| Unimed | | 689852052 | manufacture(0363-0646) |

Revised: 12/2023 Walgreen Company