# NIGHTTIME RELIEF LUBRICANT- light mineral oil, white petrolatum ointment Amerisource Bergen Drug Corp.

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## GNP Nighttime Relief Lubricant Eye Ointment 3.5g (PLD)

### Active ingredients

Light Mineral Oil 42.5%

White Petrolatum 57.3%

### Purposes

Eye Lubricant

Eye Lubricant

#### Uses

- for temporary relief of burning, irritation, and discomfort due to dryness of the eye or exposure to wind or sun
- may be used as a protectant against further irritation

### Warnings

For external use only

### Do not use

• except under the advice and supervision of a doctor

### When using this product

- replace cap after use
- to avoid contamination do not touch tip of container to any surface
- 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 (1-800-222-1222) right away.

## Directions

- pull down the lower lid of the affected eye
- apply a small amount (1/4 inch) of ointment to the inside of eyelid

## Other information

- store at 15-25°C (59-77°F)
- DO NOT FREEZE
- keep tightly closed
- do not use if it is difficult to dispense or visible particles are seen in the product
- do not roll, bend, twist, or fold tube during use
- sore away from heat

## Inactive ingredient

lanolin alcohols

## **Questions or comments?**

Call 1-888-527-4276

## **GNP Nighttime Relief Lubricant Eye Ointment 3.5g**



| NIGHTTIME RELIEF LUBRICANT<br>light mineral oil, white petrolatum ointment                             |  |                                  |       |                                 |                        |                       |                       |  |  |
|--|--|----------------------------------|-------|---------------------------------|------------------------|-----------------------|-----------------------|--|--|
| light mineral oli, w   | inite petrolat   | tum ointment                     |       |                                 |                        |                       |                       |  |  |
| Product Infor  | mation   |                                  |       |                                 |                        |                       |                       |  |  |
| Product Type   |  |                                  |       | Code (Source) NDC:46122-757     |                        |                       |                       |  |  |
|  | uct Type HUMAN OTC DRUG Item   ie of Administration OPHTHALMIC |                                  |       | Code (Source)     NDC:46122-757 |                        |                       | 0122-757              |  |  |
| Route of Admini  | stration   | OPHIHALMIC                       |       |                                 |                        |                       |                       |  |  |
|  |  |                                  |       |                                 |                        |                       |                       |  |  |
| Active Ingredient/Active Moiety  |  |                                  |       |                                 |                        |                       |                       |  |  |
| Ingredient Name  |  |                                  |       |                                 | Basis of S<br>Strength |                       | Strength              |  |  |
| <b>LIGHT MINERAL OIL</b> (UNII: N6K5787QVP) (LIGHT MINERAL OIL -<br>UNII:N6K5787QVP)                   |  |                                  |       |                                 |                        |                       | 425 mg<br>in 1 g      |  |  |
| WHITE PETROLATUM (UNII: B6E5W8RQJ4) (WHITE PETROLATUM - UNII: B6E5W8RQJ4) WHITE PETROLATUM - WHITE PET |  |                                  |       |                                 |                        | DLATUM                | 573 mg<br>in 1 g      |  |  |
|  |  |                                  |       |                                 |                        |                       |                       |  |  |
| Inactive Ingredients   |  |                                  |       |                                 |                        |                       |                       |  |  |
| Ingredient Name  |  |                                  |       |                                 |                        |                       | Strength              |  |  |
| LANOLIN ALCOHOLS (UNII: 884C3FA9HE)  |  |                                  |       |                                 |                        |                       |                       |  |  |
|  |  |                                  |       |                                 |                        |                       |                       |  |  |
| Packaging  |  |                                  |       |                                 |                        |                       |                       |  |  |
| # Item Code  | Pa   | ckage Description                |       | Marketing Start<br>Date         |                        | Marketing End<br>Date |                       |  |  |
| <b>1</b> NDC:46122-757-<br>37  | 1 in 1 BOX   |                                  | (     | 06/05/2023                      |                        |                       |                       |  |  |
| 1 3.5 g in 1 TUBE; Type 0: Not a Combination<br>Product  |  |                                  | on    |                                 |                        |                       |                       |  |  |
|  |  |                                  |       |                                 |                        |                       |                       |  |  |
| Marketing Information  |  |                                  |       |                                 |                        |                       |                       |  |  |
| Marketing<br>Category  | Applica  | tion Number or Monog<br>Citation | Iraph | Marketing Start<br>Date         |                        | Mar                   | Marketing End<br>Date |  |  |
| OTC Monograph Dru  | Ig M018  |                                  |       | 06/05/20                        | 23                     |                       |                       |  |  |
|  |  |                                  |       |                                 |                        |                       |                       |  |  |

Labeler - Amerisource Bergen Drug Corp. (007914906)

Registrant - KC Pharmaceuticals, Inc. (174450460)

| Establishment          |         |           |   |  |  |  |  |  |
|------------------------|---------|-----------|---|--|--|--|--|--|
| Name                   | Address | ID/FEI    | Business Operations   |  |  |  |  |  |
| Hanlim Pharm. Co., LTD |         | 687986034 | manufacture(46122-757) , pack(46122-757) , label(46122-757) |  |  |  |  |  |