**SODIUM CHLORIDE HYPERTONICITY- sodium chloride ointment**
**YYBA CORP**

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**Welmate ™**

**Active ingredient**

Sodium chloride 5%

**Purpose**

Hypertonicity agent

**Uses**

temporary relief of corneal edema

**Warnings**

**For external use only**

**Do not use**
except under the advice and supervision of a doctor.

**When using this product**
- it may cause temporary burning and irritation
- replace cap after use
- to avoid contamination, do not touch tip of container to any surface

**Stop use and ask a doctor if**
- condition worsens or persists for more than 72 hours
- you experience eye pain, changes in vision, continued redness or irritation of the eye

**Keep out of reach of children.**

If swallowed, get medical help or contact a Poison Control Center 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
- apply every 3 or 4 hours or as directed by a doctor

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

- **DO NOT FREEZE**
- see crimp of tube or carton for Lot Number and Expiration Date
- do not use if difficult to dispense or visible particles are seen in the product
- serious side effects associated with use of the product may be reported to the phone number provided below

**Inactive ingredients**

lanolin, mineral oil, purified water, white petrolatum

**Questions or comments?**

**Call 1-866-933-6337**

**Package/Label Principal Display Panel**


Welmate ™

NDC 73581-708-35

**EYE OINTMENT**

sodium chloride hypertonicity

ophthalmic ointment, 5%

**5%**Temporary Relief
of Corneal Edema

**STERILE**

**NET WT. 1/8 OZ. (3.5 g)**

9805700

WM15834

# SODIUM CHLORIDE HYPERTONICITY

sodium chloride ointment

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:73581-708 |
| **Route of Administration** | OPHTHALMIC | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **SODIUM CHLORIDE** (UNII: 451W47IQ8X) (SODIUM CATION - UNII:LYR4M0NH37) | SODIUM CHLORIDE | 50 mg in 1 mL |

| Inactive Ingredients | |
|---|---|
| **Ingredient Name** | **Strength** |
| **LANOLIN** (UNII: 7EV65EAW6H) | |
| **WATER** (UNII: 059QF0KO0R) | |
| **PETROLATUM** (UNII: 4T6H12BN9U) | |
| **MINERAL OIL** (UNII: T5L8T28FGP) | |

| Packaging | | | | |
|---|---|---|---|---|
| **#** | **Item Code** | **Package Description** | **Marketing Start Date** | **Marketing End Date** |
| **1** | NDC:73581-708-35 | 1 in 1 CARTON | 08/12/2024 | |
| **1** | | 3.5 mL in 1 TUBE; 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 | 08/12/2024 | |

## Labeler - YYBA CORP (006339772)

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| Bausch & Lomb Incorporated | | 079587625 | manufacture(73581-708) |

Revised: 9/2024 YYBA CORP