SILVER NITRATE- silver nitrate solution Teva Pharmaceuticals USA, Inc.

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.

Silver Nitrate Solution

Silver Nitrate Solution 0.5% 960 mL Label Text

NDC 0093-9614-13

Silver Nitrate

Solution

0.5%

FOR EXTERNAL USE ONLY.

Rx only

960 mL (32 fl oz)

TEVA

DESCRIPTION: Silver nitrate solution is a 0.5% solution of

silver nitrate in a water medium. It is a topical anti-infective.

WARNINGS: When ingested, silver nitrate is highly toxic to

the gastrointestinal tract and central nervous system.

Swallowing can cause severe gastroenteritis that may end

fatally. Sodium chloride may be used by gastric lavage to

remove the chemical. Caustic and irritating to the skin and

mucous membranes.

PRECAUTIONS: Silver nitrate solution must be handled carefully,

since it tends to stain the skin, utensils, clothing and linens.

DOSAGE: Topical as directed by the physician.

Store at 20° to 25°C (68° to 77°F) [See USP Controlled

Room Temperature]. Do not freeze. Protect from light.

KEEP THIS AND ALL MEDICATIONS OUT OF THE REACH

OF CHILDREN.

Manufactured By:

HALO PHARMACEUTICALS

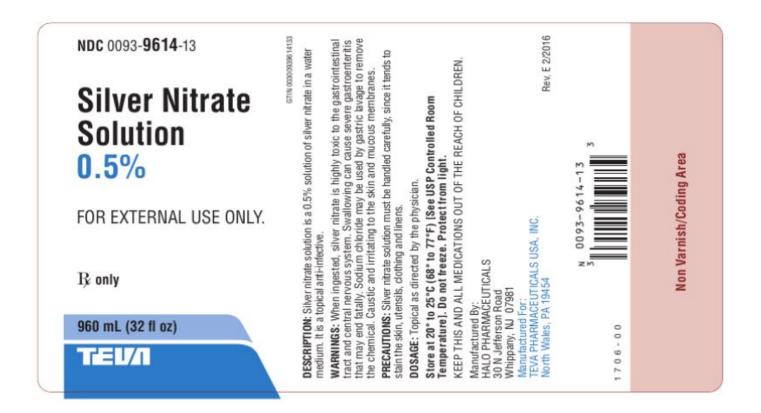
30 N Jefferson Road

Whippany, NJ 07981

Manufactured For:

TEVA PHARMACEUTICALS USA, INC.

Package/Label Display Panel



SILVER NITRATE

silver nitrate solution

| Product Information | | | | | |
|-------------------------|-------------------------|--------------------|---------------|--|--|
| Product Type | HUMAN PRESCRIPTION DRUG | Item Code (Source) | NDC:0093-9614 | | |
| Route of Administration | TOPICAL | | | | |

| Active Ingredient/Active Moiety | | |
|---|-------------------|-----------------|
| Ingredient Name | Basis of Strength | Strength |
| SILVER NITRATE (UNII: 95IT3W8JZE) (SILVER CATION - UNII:57N7B0K90A) | SILVER NITRATE | 0.005 g in 1 mL |

| Inactive Ingredients | | | | | |
|--------------------------|----------|--|--|--|--|
| Ingredient Name | Strength | | | | |
| WATER (UNII: 059QF0KO0R) | | | | | |
| | | | | | |

| Packaging | | | | | | | | |
|-----------------------|------------------|--------|--|-----------------------------|---------------------------|--|--|--|
| # | Item Code | | Package Description | Marketing Start Date | Marketing End Date | | | |
| 1 | NDC:0093-9614-13 | 960 mL | in 1 BOTTLE; Type 0: Not a Combination Product | 0 1/0 1/19 9 1 | | | | |
| | | | | | | | | |
| | | | | | | | | |
| Marketing Information | | | | | | | | |
| | Marketing Categ | ory | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date | | | |
| Ul | NAPPRO VED DRUG | OTHER | | 0 1/0 1/19 9 1 | | | | |
| | | | | | | | | |

Labeler - Teva Pharmaceuticals USA, Inc. (001627975)

Revised: 2/2016 Teva Pharmaceuticals USA, Inc.