

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.

North Wales, PA 19454

333-32-101019 Rev. E 2/2016

102121 0786-01

Package/Label Display Panel

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1706-00

Non Varnish/Coding Area

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	01/01/1991	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
UNAPPROVED DRUG OTHER		01/01/1991	

Labeler - Teva Pharmaceuticals USA, Inc. (001627975)

Revised: 2/2016

Teva Pharmaceuticals USA, Inc.