TINCTURE MERTHIOLATE- benzalkonium chloride tincture DLC Laboratories, Inc

De La Cruz Tincture Merthiolate

Active Ingredient

Benzalkonium chloride 0.13%

Purpose

First Aid Antiseptic

Uses

First aid antiseptic to help prevent infection in minor:

cuts - scrapes - burns

Warnings

For external use only.

Flammable. Keep away from sparks, heat and fire.

Consult a doctor before use if you have - deep or puncture wounds - animal bites - serious burns

When using this product

- do not use in or near the eyes
- do not apply in large quantities or over large areas of the body
- do not apply over raw surtaces or blistered areas
- do not use longer than 1 week unless directed by a doctor

Stop use and consult a doctor if

• the condition persists or gets worse

Keep out of reach of children.

II swallowed, get medical help or contact a Poison Control Center immediately.

Directions

- adults and children 2 years of age and older: clean the affected area
- apply a small amount of this product to the area 1 to 3 times daily _ may be covered with a sterile bandage. If bandaged, let dry first
- children under 2 years of age: ask a doctor

Other Information

will stain skin and clothing

Inactive Ingredients

acetone, alcohol, DandC red no. 22, purified water

Product Labeling





Tincture Merthiolate
48% Alcohol
Mercury Free
First Aid Antiseptic
For External Use Only
1 FL OZ (30 mL)
Distributed by

De La Cruz Products

A Division of DLC Laboratories, Inc.

Paramount, CA 90723 USA

Questions: 1-800-858-3889

www.dlclabs.com

Warnings: FLAMMABLE, KEEP AWAY FROM SPARKS, HEAT AND FIRE.

TINCTURE MERTHIOLATE

benzalkonium chloride tincture

Product Type HUMAN OTC DRUG Item Code (Source) NDC:24286-1532

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength

BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - BENZALKONIUM - UNII:7N6JUD5X6Y)

BENZALKONIUM - BENZALKONIUM - CHLORIDE in 1 mL

Inactive Ingredients

Ingredient Name	Strength			
ACETONE (UNII: 1364PS73AF)				
ALCOHOL (UNII: 3K9958V90M)				
D&C RED NO. 22 (UNII: 1678RKX8RT)				

WATER (UNII: 059QF0KO0R)

Packaging

# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:24286- 1532-7	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/13/2017	

Marketing Information

Marketing Applicatio Category	n Number or Monograph	Marketing Start	Marketing End
	Citation	Date	Date
OTC Monograph Drug M003		03/22/2013	

Registrant - Pharma Nobis, LLC (118564114)

Establishment				
Name	Address	ID/FEI	Business Operations	
Pharma Nobis, LLC		118564114	analysis(24286-1532), manufacture(24286-1532), pack(24286-1532), label(24286-1532)	

Revised: 12/2023 DLC Laboratories, Inc