# TINCTURE MERTHIOLATE- benzalkonium chloride tincture DLC Laboratories, Inc

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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#### De La Cruz Tincture Merthiolate

**Drug Facts** 

### **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 eves
- do not apply over large areas of the body
- do not apply over raw surfaces 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.

If 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, D&C red no. 22, purified water

## Questions

1-800-858-3889

Distributed by: De La Cruz Products A Division of DLC Laboratories, Inc. Paramount, CA 90723 USA

#### PRINCIPAL DISPLAY PANEL - 30 mL Bottle Label

De La Cruz®

Tincture

Merthiolate

48%

Alcohol

Mercury-Free

First Aid Antiseptic

TRUSTED QUALITY

For External Use Only

1 FL OZ (30 mL)

Datos Medicinales



Questions / Preguntas: 1-800-858-3889 A Division of DLC Laboratories, Inc. Distributed by / Distribuido por: © 2013 DL Paramount, CA 90723 USA De La Cruz Products www.dlclabs.com

P0947-GZT

**ADVERTENCIA:** INFLAMABLE. MANTÉNGASE ALEJADO DE CHISPAS, CALOR Y FUEGO. WARNING: FLAMMABLE. KEEP AWAY FROM SPARKS, HEAT AND FIRE

## Drug Facts Active ingredient

Purpose First aid

Benzalkonium chloride 0.13%......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 over large areas of the body do not apply over raw surfaces 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. If 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

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#### benzalkonium chloride tincture

### **Product Information**

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
<b>BENZALKO NIUM CHLO RIDE</b> (UNII: F5UM2KM3W7) (BENZALKO NIUM - UNII:7N6 JUD5X6 Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
ACETONE (UNII: 1364PS73AF)	
ALCOHOL (UNII: 3K9958V90M)	
<b>D&amp;C RED NO. 22</b> (UNII: 1678 RKX8 RT)	
WATER (UNII: 059QF0KO0R)	

ı	Packaging			
ı	# Item Code	Package Description	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
ı	1 NDC:24286-1532-7	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/13/20 17	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/22/2013	

## Labeler - DLC Laboratories, Inc (093351930)

## Registrant - Humco Holding group, Inc. (825672884)

Revised: 12/2019 DLC Laboratories, Inc