

DDM MERTHIOLATE- benzalkonium chloride liquid

Discount Drug Mart

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.

DDM Merthiolate

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Drug Facts

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antiseptic

Indications

First aid antiseptic to help skin infection in minor: cuts, scrapes, burns, insect bites.

Warnings

For external use only.

Ask a doctor before use if you have

Deep or puncture wounds

Animal bites

Serious burns

When using this product do not

Get into the eyes

Apply over large areas of the body

Apply over raw surfaces or blistered areas

Use longer than one week unless directed by doctor.

Stop use and ask a doctor if

Redness, swelling or pain persists or increases.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center immediately. Soap will deactivate the effects of this product.

Directions

- Adults and children 2 years and older. Clean the affected area; apply a small amount on the area 1 to 3 times daily; may be covered with a sterile bandage. If bandaged, let dry first.
- Children under 2 yrs. of age: Consult a doctor.

Inactive Ingredient

Acetone, FD&C Red No 4. Purified water.

Label

<p>Drug Facts</p> <table border="1"> <tr> <td>Active ingredient</td> <td>Purpose</td> </tr> <tr> <td>Benzalkonium chloride 0.13%.....</td> <td>Antiseptic</td> </tr> </table> <p>Uses First aid antiseptic to help prevent skin infection in minor ■ cuts ■ scrapes ■ burns</p> <p>Warnings For external use only Ask a doctor before use if you have ■ deep or puncture wounds ■ animal bites ■ serious burns</p> <p>When using this product do not ■ get into the eyes ■ apply over large areas of the body ■ over raw surfaces or blistered areas ■ use longer than one week unless directed by a doctor</p> <p>Stop use and consult a doctor if the condition persists or gets worse. Continued ▶</p> <p style="text-align: right;">MADE IN USA</p>	Active ingredient	Purpose	Benzalkonium chloride 0.13%.....	Antiseptic		<p>(Drug Facts continued)</p> <p>Do not use longer than 1 week unless directed by a doctor</p> <p>Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center immediately. Soap will deactivate the effects of this product.</p> <p>Directions ■ Clean the affected area apply a small amount on the area 1 to 3 times daily may be covered with a sterile bandage. If bandaged, let dry first.</p> <p>Inactive ingredients Alcohol 10%, Acetone, FD&C Red No. 4, Purified water.</p>	 <p>0 93351 01326 3</p>
Active ingredient	Purpose						
Benzalkonium chloride 0.13%.....	Antiseptic						
<p>MERTHIOLATE Mercury-Free Formula! First Aid Antiseptic</p>		<p>Helps prevent infection in minor cuts, scrapes & burns</p> <p>2 FL OZ (59mL)</p>					
<p>DISTRIBUTED BY: R051413.RLG DRUG MART-FOOD FAIR 211 Commerce Dr. MEDINA, OHIO Zip 44256 Questions or Comments? 1-800-833-6278 www.DISCOUNT-DRUGMART.COM</p>							

DDM MERTHIOLATE

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53943-749
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ACETONE (UNII: 1364PS73AF)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53943-749-92	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/26/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/26/2017	

Labeler - Discount Drug Mart (047741335)

Registrant - Humco Holding Group, Inc. (825672884)

Establishment

Name	Address	ID/FEI	Business Operations
Humco Holding Group, Inc.		825672884	manufacture(53943-749) , analysis(53943-749) , pack(53943-749) , label(53943-749)

Revised: 11/2017

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