

**PREMIER VALUE MERTHIOALATE- benzalkonium chloride liquid**  
**Chain Drug Consortium**

*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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**Premier Value 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><b>Drug Facts</b></p> <table border="1"> <tr> <th style="text-align: left;">Active ingredient</th> <th style="text-align: left;">Purpose</th> </tr> <tr> <td>Benzalkonium chloride 0.13%</td> <td>Antiseptic</td> </tr> </table> <p><b>Uses</b> First aid antiseptic to help prevent skin infection in minor ■ cuts ■ scrapes ■ burns</p> <p><b>Warnings</b> 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.</p> <p style="text-align: right;"><b>Continued</b> ▶</p>	Active ingredient	Purpose	Benzalkonium chloride 0.13%	Antiseptic	 <p>NDC 68016-437-00</p> <p><b>Antiseptic</b> <b>MERTHIOALATE</b> <b>Mercury-Free Formula</b></p> <p>• Helps prevent infection in minor cuts, scrapes and burns</p> <p>NET WT. 2 FL. OZ (59 mL)</p> <p>MADE IN USA</p>	<p><b>(Drug Facts continued)</b></p> <p>Do not use longer than 1 week unless directed by a doctor</p> <p><b>Keep out of reach of children.</b> If swallowed, get medical help or contact a Poison Control Center immediately. Soap will deactivate the effects of this product.</p> <p><b>Directions</b> ■ 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><b>Inactive ingredients</b> Alcohol 10%, Acetone, FD&amp;C Red No. 4, Purified water.</p> <p style="text-align: right;">R120313RLG</p> <p>DISTRIBUTED BY: CHAIN DRUG CONSORTIUM 3301 NW BOCA RATON BLVD SUITE 101, BOCA RATON, FL 33431 MADE IN U.S.A.</p>
Active ingredient	Purpose					
Benzalkonium chloride 0.13%	Antiseptic					

## PREMIER VALUE MERTHIOALATE

benzalkonium chloride liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:68016-437
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

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

### Inactive Ingredients

Ingredient Name	Strength
<b>ACETONE</b> (UNII: 1364PS73AF)	
<b>FD&amp;C RED NO. 4</b> (UNII: X3W0AM1JLX)	
<b>WATER</b> (UNII: 059QF0KO0R)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68016-437-00	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** - Chain Drug Consortium (101668460)

**Registrant** - Humco Holding Group, Inc. (825672884)

## Establishment

Name	Address	ID/FEI	Business Operations
Humco Holding Group, Inc.		825672884	manufacture(68016-437) , analysis(68016-437) , pack(68016-437) , label(68016-437)

Revised: 12/2017

Chain Drug Consortium