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.

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

PREMIER VALUE MERTHIOALATE

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68016-437
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: 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