CVS MERTHIOLATE- benzalkonium chloride liquid
CVS Pharmacy

CVS 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

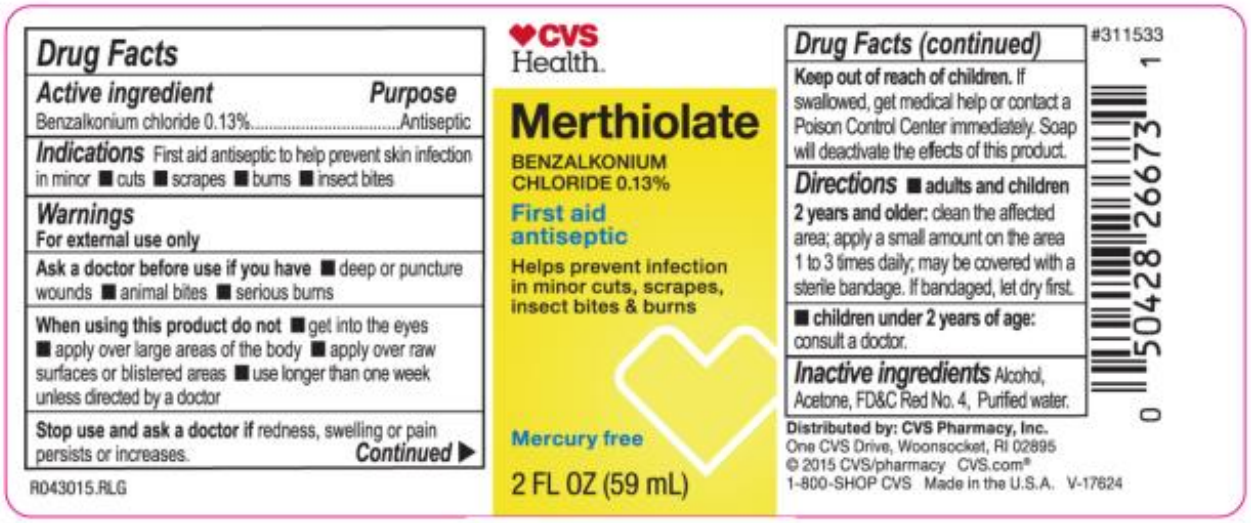
- **Adults and children 2 years and older**. Clean the affected area; apply a small amount on the area 1 to 3 times daily; may be covered with a sterile bandage. If bandaged, let dry first.
- **Children under 2 yrs. of age:** Consult a doctor.

Inactive Ingredient

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

PRINCIPAL DISPLAY PANEL

CVS
Merthiolate
Benzalkonium
Chloride 0.13%
Mercury free
2 FL OZ (59 mL)



CVS MERTHIOLATE

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69842-749
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: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69842-749-92	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/2008	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	01/01/2008	

Labeler - CVS Pharmacy (062312574)

Registrant - Pharma Nobis, LLC (118564114)

Establishment

Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	analysis(69842-749) , manufacture(69842-749) , pack(69842-749) , label(69842-749)

Revised: 12/2023 CVS Pharmacy