QUALITY CHOICE MERTHIOLATE- benzalkonium chloride liquid
Chain Drug Market Association

Quality Choice Merthiolate

Drug Facts

Active Ingredient

Benzalkonium chloride 0.13%

Purpose

Antiseptic

Uses

first aid to help prevent skin infection in minor cuts, scrapes, burns and 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 eyes. If contact occurs, rinse eyes throughly with water. do not apply over large areas of the body. do not use over raw surfaces or blistered areas. do not use longer than 1 week unless directed by a doctor.

Stop use and ask a doctor if

condition persists or gets worse. symptoms clear up and occur again within a few days.

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center immediately.

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 it dry first. children under 2 years of age, do not use, consult a doctor.

Inactive ingredients

Alcohol 10%, Acetone, FD&C Red No 4, purified water

Principal display panel

Label

QUALITY CHOICE MERTHIOLATE

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63868-493
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**
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	

ACETONE (UNII: 1364PS73AF)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-493-02	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/12/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	03/25/1998	

Labeler - Chain Drug Market Association (011920774)

Registrant - Pharma Nobis, LLC (118564114)

Establishment

Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	analysis(63868-493) , manufacture(63868-493) , pack(63868-493) , label(63868-493)

Revised: 12/2023 Chain Drug Market Association