STERILE ANTIBACTERIAL WOUND WASH - benzalkonium chloride spray
Ascot Pharmaceuticals, Inc.

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.

Drug Facts

Active ingredients

Benzalkonium Chloride 0.13%

Purpose

First Aid Antiseptic

Uses

first aid to help prevent the risk of bacterial contamination in minor cuts, scrapes and burns.

Warnings

For external use only

Do not use

- in eyes
- over large areas of the body
- longer than 1 week unless directed by a doctor

Ask a doctor before use

in the event of deep puncture wounds, animal bites or severe burns

When using this product

use only as directed

Stop use and ask a doctor if

- conditions worsens
- symptoms persists for more than seven days
- symptoms disappear and reappear again within a few days

Keep out of reach of childrens

If swallowed get medical help or contact a Poison Control Center right away

Directions

- clean the affected area
- shake well
- remove seal and cap
- spray a small amount of this product on the area 1 to 3 times a day
- may be covered with a sterile bandage. If bandaged, let dry first

Other information

- contents under pressure
- do not puncture or incinerate can
- do not drop can
- do not store at temperatures above 120 degrees F.
- do not use if tamper evident seal is broken

Inactive ingredients

Sodium Chloride U.S.P., Aloe Barbadensis Leaf Extract , Purified Water

Questions?

1-800-831-7210

Principal Display Panel

NDC 70043-390-04

Ascot Pharmaceutical Inc.

Sterile Antibacterial

WOUND WASH

FIRST AID CLEANSER
- Cleans wounds painlessly and effectively
- No burn or sting
- No Preservatives
- No alcohol
- No fragrance

ECO-FRIENDLY

No Fluorinated Hydrocarbon Propellants

- COMPRESSED AIR IS THE PROPELLANT
- SPRAYS UPSIDE DOWN OR AT ANY ANGLE
- NO PUMPING NEEDED

360^0 CONTINUOUS SPRAY

4 FL.OZ (120 mL)

STERILE ANTIBACTERIAL WOUND WASH

benzalkonium chloride spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70043-390
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**
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70043-390-04	120 mL in 1 CAN; Type 0: Not a Combination Product	06/23/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/23/2016	

Labeler - Ascot Pharmaceuticals, Inc. (079915219)

Establishment

Name	Address	ID/FEI	Business Operations
BOV Solutions Inc		800463486	manufacture(70043-390)

Revised: 6/2016 Ascot Pharmaceuticals, Inc.