

**BAND AID ANTISEPTIC CLEANSING- benzalkonium chloride aerosol, foam
Johnson & Johnson Consumer 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.

Band Aid Antiseptic Cleansing

Drug Facts

Active ingredient

Benzalkonium Cl (0.13%)

Purposes

First aid antiseptic

Use

first aid to help prevent infection in minor: · cuts · scrapes · burns

Warnings

- **For external use only.**

Do not use · in the eyes · over large area of the body · longer than 1 week

- **Ask a doctor before use if you have · deep or puncture wounds · animal bites · serious burns**

- **Stop use and ask a doctor if the condition persists or gets worse**

Keep out of reach of children.

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

Directions

adults and children 2 years of age and older:

clean the affected area

apply a small amount of this product on the area 1 to 3 times daily

may be covered with a sterile bandage

if bandaged, let dry first

children under 2 years of age: consult a doctor

Other information

protect from freezing

Inactive ingredients

Water, Poloxamer 188, Sodium Chloride, Disodium EDTA, Sodium Hydroxide, Aloe Barbados Leaf Juice, Citric Acid

Questions?

call 800-526-3967 or 215-273-8755 (collect).

Distributed by:

JOHNSON & JOHNSON CONSUMER INC.

Skillman, NJ 08558

PRINCIPAL DISPLAY PANEL - 68 mL Bottle

BAND-AID®

BRAND OF FIRST AID PRODUCTS

Johnson & Johnson

ANTISEPTIC

CLEANISING

FOAM

FIRST AID ANTIBIOTIC

Kids

Kills 99.99% of germs*

to prevent infection

No sting

Contains aloe

For minor cuts,

scrapes & burns

2.3 FL OZ (68mL)



BAND AID ANTISEPTIC CLEANSING

benzalkonium chloride aerosol, foam

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69968-0728
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)	
POLOXAMER 188 (UNII: LQA7B6G8JG)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0728-1	68 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	12/29/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	12/29/2021	

Labeler - Johnson & Johnson Consumer Inc. (118772437)

Revised: 1/2023

Johnson & Johnson Consumer Inc.