NEOSPORIN FIRST AID ANTISEPTIC FOAMING FOR KIDS- benzalkonium chloride liquid 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.

NEOSPORIN[®] FIRST AID ANTISEPTIC FOAMING LIQUID FOR KIDS[®]

Drug Facts

Active ingredient

Benzalkonium Cl 0.13%

Purpose

First aid antiseptic

Uses

first aid to help prevent infection in minor:

- cuts
- scrapes
- burns

Warnings

For external use only

Do not use

- in the eyes
- over large areas 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

- store at 20° to 25°C (68° to 77°F)
- protect from freezing

Inactive ingredients

aloe barbadensis leaf juice, citric acid, disodium EDTA, poloxamer 188, sodium chloride, sodium hydroxide, water

Questions?

call toll-free 800-223-0182 or 215-273-8755 (collect)

Distributed by: JOHNSON & JOHNSON CONSUMER INC. Skillman, NJ 08558

PRINCIPAL DISPLAY PANEL - 68 mL Bottle Label

See New Directions!

FROM THE #1 DOCTOR RECOMMENDED BRAND

NEOSPORIN[®] BENZALKONIUM CHLORIDE FIRST AID ANTISEPTIC FOAMING LIQUID

FOR KIDS Ages 2 and up

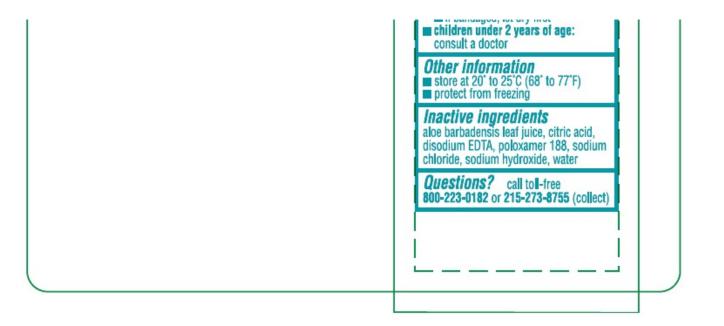
No Sting

Kills Germs to Help Prevent Infection

2.3 fl oz (68 mL)







NEOSPORIN FIRST AID ANTISEPTIC FOAMING FOR KIDS benzalkonium chloride liquid **Product Information Product Type** HUMAN OTC DRUG Item Code (Source) NDC:69968-0220 **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) Citric Acid Monohydrate (UNII: 2968PHW8QP) Edetate Disodium Anhydrous (UNII: 8NLQ36F6MM) Poloxamer 188 (UNII: LQA7B6G8JG) Sodium Chloride (UNII: 451W47IQ8X) Sodium Hydroxide (UNII: 55X04QC32I) Water (UNII: 059QF0KO0R) Packaging **Marketing Start Marketing End** # Item Code **Package Description** Date Date 1 NDC:69968-0220- 68 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product 06/01/2009 Product

Marketing Information		
Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
part333A	06/01/2009	
1	Application Number or Monograph Citation	Application Number or Monograph Citation Marketing Start Date

Labeler - Johnson & Johnson Consumer Inc. (002347102)

Revised: 4/2020

Johnson & Johnson Consumer Inc.