HILLYARD ALCOHOL FREE FOAMING INSTANT HAND SANITIZER- benzalkonium chloride liquid
Hillyard GMP

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.

Hillyard Alcohol Free Foaming Instant Hand Sanitizer

Active ingredient

Benzalkonium Chloride 0.10%

Purpose

Antimicrobial

Uses

- For hand sanitizing to decrease bacteria on the skin
- Recommended for repeated use.

Warnings

- For External Use Only

When using this product

avoid contact with eyes. In case of eye contact, flush eyes thoroughly with water.

Stop use and ask a doctor if

irritation or redness develops, or if condition persists for more than 72 hours.

Keep Out of Reach of Children

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

Directions

- Pump small amount of foam into palm of hand
- Rub thoroughly over all surfaces of both hands
- Rub hands together briskly until dry

Inactive Ingredients

Water, Cetrimonium Chloride, Laurtrimonium Chloride, Dihydroxyethyl Cocamine Oxide, Glycereth-17 Cocoate, Phenoxyethanol and Citric Acid.

Package/Label Principal Display Panel

Hillyard
Alcohol Free Foaming
Instant Hand Sanitizer

HIL0040303

affinity
DISPENSING SYSTEM

Place cartridge into dispenser, label facing back, and gently push pump into receiver until it clicks into place.

To remove the cartridge and pump, rotate the green ring clockwise and pull the pump out.

HILLYARD INDUSTRIES
PO Box 909,
St. Joseph, MO
64502-0909 U.S.A.
Telephone: 816-233-1321
www.hillyard.com

NET CONTENTS 1.25 L (42.3 FL. OZ.)
PEEL HERE

Drug Facts

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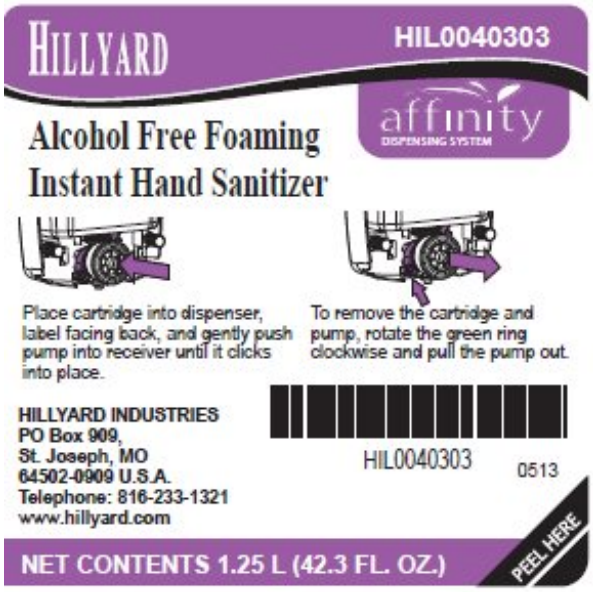
NDC 76402-403-03

HIL0040303



Carton Label

HILLYARD ALCOHOL FREE FOAMING INSTANT HAND SANITIZER

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:76402-403
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.1 g in 100 mL

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
LAURTRIMONIUM CHLORIDE (UNII: A81MSI0FIC)	
DIHYDROXYETHYL COCAMINE OXIDE (UNII: 8AR51R3BL5)	
GLYCERETH-17 COCOATE (UNII: 3057VPT0KC)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76402-403-06	3785 mL in 1 JUG; Type 0: Not a Combination Product	02/29/2012	
2	NDC:76402-403-03	1250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/02/2012	
3	NDC:76402-403-02	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/04/2012	
4	NDC:76402-403-82	414 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/18/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	02/29/2012	

Labeler - Hillyard GMP (969081483)

Establishment

Name	Address	ID/FEI	Business Operations
Hillyard GMP		969081483	MANUFACTURE(76402-403)

Revised: 1/2015 Hillyard GMP