

OPI PERSONAL CARE SUPPLIES INSTANT FOAMING HAND SANITIZER NON ALCOHOL- husky 514 solution

OPI Correctional Industries

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.

Instant Foaming Hand Sanitizer Non Alcohol

Drug Facts

Active Ingredient

Benzalkonium Chloride 0.1%

Purpose

Antimicrobial

Uses

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

Warning

For external use only

When using this product avoid contact with eyes. In case of eye contact, flush eyes 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 a 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, dihydroxypropyl PEG-5 linoleammonium chloride, glycereth-2 cocoate, behentrimonium chloride, dihydroxyethyl cocamine oxide, fragrance

Principal Display Panel

OPI Personal Care Supplies

Instant Foaming Hand Sanitizer Non Alcohol

Ready-to-Use

- Enhanced with Moisturizers

- Kills disease causing germs within seconds
- Effective against MRSA, VRE, E. coli (0157:H7) Staphylococcus, Streptococcus and other organisms
- Assists with OSHA Bloodborne Pathogen Standard Compliance

See Drug Facts panel for additional information.

For Hospital and Professional Use Only

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514-001-093-501
Reorder 22021
Sold By:



This container is coded for Recycling

OPI CORRECTIONAL INDUSTRIES
Columbus, OH 43222
800.237.3454 • www.opi.ohio.gov



INSTANT FOAMING HAND SANITIZER

NON-ALCOHOL READY-TO-USE

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Net Contents: One Gallon (128 Fl. Oz.) 3.78 liters

OPI PERSONAL CARE SUPPLIES INSTANT FOAMING HAND SANITIZER NON ALCOHOL

husky 514 solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:43196-514
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
benzalkonium chloride (UNII: F5UM2KM3W7) (benzalkonium - UNII:7N6JUD5X6Y)	benzalkonium chloride	1 g in 1000 mL

Inactive Ingredients

Ingredient Name	Strength
water (UNII: 059QF0K00R)	
dihydroxypropyl peg-5 linoleammonium chloride (UNII: 0 Y0 NQR2GH1)	
glycereth-2 cocoate (UNII: JWM00VS7HC)	
behentrimonium chloride (UNII: X7GNG3S47T)	
dihydroxyethyl cocamine oxide (UNII: 8 AR51R3BL5)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:43196-514-05	3785 mL in 1 BOTTLE		
2	NDC:43196-514-83	550 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333	11/16/2009	

Labeler - OPI Correctional Industries (809174501)

Establishment

Name	Address	ID/FEI	Business Operations
Canberra Corporation		068080621	MANUFACTURE

Revised: 3/2010

OPI Correctional Industries