FIKES NORTHWEST NON-ALCOHOL FOAMING INSTANT HAND SANITIZER- hus ky 514 solution

Fikes Northwest, Corp.

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.

Non AlcoholFoaming Instant Hand Sanitizer

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, dihydroxpropyl PEG-5 linoleammonium chloride, glycereth-2 cocoate, behentrimonium chloride, dihydroxyethyl cocamine oxide, fragrance

Principal Display Panel

Fikes Northwest

Non AlcoholFoaming Instant Hand Sanitizer

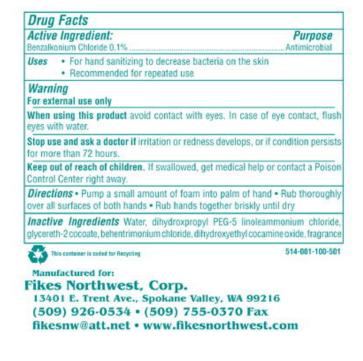
- Enhanced with Moisturizers
- Kills disease causing germs within seconds
- Effective against MRSA, VRE, E. coli (0157:H7) Staphylococcus, Streptococcus and other organisms

• Assists with OSHA Bloodborne Pathogen Standard Compliance

See side panel for additional information.

For Hospital and Professional Use Only

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





FIKES NORTHWEST NON-ALCOHOL FOAMING INSTANT HAND SANITIZER

husky 514 solution

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50036-514
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: 059QF0KO0R)	
dihydroxypropyl peg-5 linoleammonium chloride (UNII: 0 Y0 NQR2GH1)	
glycereth-2 cocoate (UNII: JWM00VS7HC)	
behentrimonium chloride (UNII: X7GNG3S47T)	
dihydroxyethyl cocamine oxide (UNII: 8AR51R3BL5)	

Packaging					
# Item Code	Package Description	Marketing	Start Date	Marketing End Date	
1 NDC:50036-514-05	3785 mL in 1 BOTTLE				
Mr. J. C. at T. C. at a control of					
Marketing Information					
Manlarda Catagoni	Application Number or Monog	wanh Citation	Marketing Start	Date Marketing End Date	
Marketing Category	Application Number of Monog	rapii Citation	Marketing Start	Date Marketing End Date	
OTC monograph not final	part333	rapii Citation	11/16/2009	Date Marketing Lift Date	

Labeler - Fikes Northwest, Corp. (167376284)

Registrant - Fikes Northwest, Corp. (167376284)

Establishment					
Name	Address	ID/FEI	Business Operations		
Canberra Corporation		068080621	MANUFACTURE		

Revised: 1/2010 Fikes Northwest, Corp.