

**FIKES NORTHWEST NON-ALCOHOL FOAMING INSTANT HAND SANITIZER- husky 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.*

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**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, dihydroxypropyl 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

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 This container is coded for Recycling	514-001-100-501
Manufactured for: <b>Fikes Northwest, Corp.</b> 13401 E. Trent Ave., Spokane Valley, WA 99216 (509) 926-0534 • (509) 755-0370 Fax fikesnw@att.net • www.fikesnorthwest.com	

# Fikes<sup>northwest</sup>

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## FIKES NORTHWEST NON-ALCOHOL FOAMING INSTANT HAND SANITIZER

husky 514 solution

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:50036-514
<b>Route of Administration</b>	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: 0Y0NQR2GH1)	
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		

**Marketing Information**

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

**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.