

**HUSKY 514 NON-ALCOHOL FOAMING INSTANT HAND SANITIZER- husky 514 solution
Canberra Corporation**

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.

Husky® 514

NON-ALCOHOL FOAMING 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 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

Manufactured By:

CANBERRA CORPORATION

3610 Holland-Sylvania Rd.

Toledo, Ohio 43615

Label Content for 128 Fl. Oz:

Husky®

NON-ALCOHOL FOAMING INSTANT HAND SANITIZER

- Enhanced with Moisturizers
- Kills disease causing germs within seconds
- Effective against MRSA, VRE, E. coli (O157: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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FOAMING SOAPS
SKIN CARE • SKIN CARE

Label Content for 18.60 Fl. Oz:

Husky®

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Net Contents: 18.60 fl. oz./550 mL.



Label Content for 250 mL:

Husky

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Net Contents: 250 mL

514-001-001-S01F



Label Content for 1.69 Fl Oz:

Husky

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Net Contents: 1.69 fl. Oz (50 mL)



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husky 514 solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63779-514
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: 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:63779-514-10	3785 mL in 1 BOTTLE		
2	NDC:63779-514-82	550 mL in 1 BOTTLE		
3	NDC:63779-514-99	250 mL in 1 BOTTLE		
4	NDC:63779-514-42	50 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 - Canberra Corporation (068080621)

Registrant - Canberra Corporation (068080621)

Establishment

Name	Address	ID/FEI	Business Operations
Canberra Corporation		068080621	MANUFACTURE

Revised: 11/2009

Canberra Corporation