# HUSKY 515 INSTANT HAND SANITIZER - ethanol gel 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.

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# **Drug Facts**

# **Active Ingredient:**

Ethanol (60% v/v)

#### Purpose:

Anti-Microbial Hand Sanitizer

#### Uses

- Helps reduce bacteria that potentially can cause disease
- Helps prevent cross contamination by hand contact
- Recommended for repeated use

#### **Warnings**

- For external use only
- Flammable, keep away from fire, heat, or flame
- Keep out of reach of children.

#### Do not use near eyes

- In case of eye contact flush with water for 15 minutes
- If irritation persists stop use of product and get medical attention
- In case of accidental ingestion seek medical attention or contact a poison control center immediately.

#### **Directions**

- Use no water or towels
- Apply appropriate amount of product to palm of hand
- Rub until hands are completely covered
- Agitate lightly until dry
- Let air dry for 15 seconds
- Do not rinse or wipe with towel.

#### Other Information

• Store in a cool dry place below 104° F.

#### **Inactive Ingredients**

Water, Carbomer, Triethanolamine, PEG-75 Lanolin, Aloe Vera Gel, Fragrance.

#### **Principal Display Panel**

#### **Bottle Label**

## **HUSKY**®

515

**INSTANT HAND** 

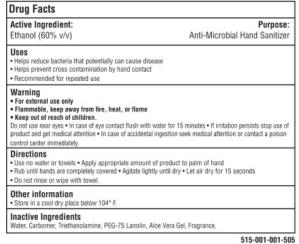
#### SANITIZER

(GEL-TYPE)

- 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

# For Hospital and Professional Use Only

### See Drug Facts panel for additional information.



515-001-001



Manufactured By: CANBERRA CORPORATION 3610 Holland-Sylvania Rd. Toledo, Ohio 43615





#### **HUSKY 515 INSTANT HAND SANITIZER**

ethanol gel

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63779-515	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
alcohol (UNII: 3K9958V90M) (alcohol - UNII:3K9958V90M)	alcohol	600 mL in 1 L		

#### **Inactive Ingredients**

Ingredient Name	Strength
water (UNII: 059QF0KO0R)	
carbomer homopolymer type c (UNII: 4Q93RCW27E)	
aloe (UNII: V5VD430 YW9)	
trolamine (UNII: 9O3K93S3TK)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63779-515-05	3.785 L in 1 BOTTLE		
2	NDC:63779-515-15	0.044 L in 1 BOTTLE		
3	NDC:63779-515-27	0.118 L in 1 BOTTLE		
4	NDC:63779-515-62	0.237 L in 1 BOTTLE		
5	NDC:63779-515-80	0.800 L in 1 BAG		
6	NDC:63779-515-88	0.237 L in 1 BOTTLE		
7	NDC:63779-515-89	0.490 L in 1 BOTTLE		
8	NDC:63779-515-20	1.100 L in 1 BOTTLE		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	10/28/1998		

# Labeler - Canberra Corporation (068080621)

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

Revised: 8/2011 Canberra Corporation