## HUSKY 575 FOAMING ANTISEPTIC HAND CLEANER- triclos an liquid 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:**

Triclosan (0.35%)

#### **Purpose:**

Antiseptic Hand Cleanser

#### Uses

- Handwash to help reduce bacteria that potentially can cause disease.
- Recommended for repeated use.

#### **Warnings**

- For external use only
- Do not use near eyes
- Keep out of reach of children.

Discontinue use if irritation and redness develop. If condition persists more than 72 hours, consult a doctor.

In case of eye contact flush with water for 15 minutes. If irritation persists, get medical attention

• In case of accidental ingestion, seek medical attention or contact a poison control center immediately.

#### Directions

- Wet hands and forearms
- Apply appropriate amount (palmful) to hands and forearms
- Scrub hands and forearms thoroughly for 60 seconds
- Use scrub and/or nail brush to aid in cleaning.
- Rinse well
- Wipe dry with towel, repeat if necessary.

#### Other Information

Assists with OSHA bloodborne pathogen standard compliance

#### **Inactive Ingredients**

Water, Sodium Laureth Sulfate, Sodium Lauryl Sulfate, Propylene Glycol, Lauramide DEA, Cocamide DEA, Cocamidopropyl betaine, Sodium Cumene Sulfonate, Citric Acid, Fragrance, DMDM Hydantoin, Sodium Chloride, FD&C Green #3.

#### **Principal Display Panel**

#### **Bottle Label:**

#### **HUSKY**®

575

#### FOAMING ANTISEPTIC HAND CLEANER

A Powerful Antiseptic Cleanser for Handwashing

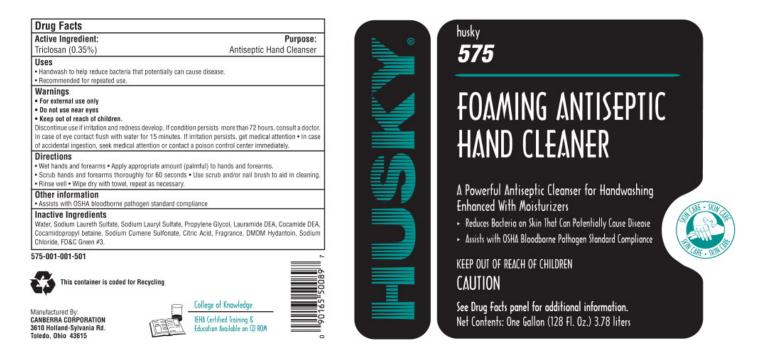
**Enhanced** with Moisturizers

- Reduces Bacteria on Skin That Can Potentially Cause Disease
- Assists with OSHA Bloodborne Pathogen Standard Compliance

#### KEEP OUT OF REACH OF CHILDREN

#### **CAUTION**

See Drug Facts panel for additional information.



# HUSKY 575 FOAMING ANTISEPTIC HAND CLEANER triclosan liquid

**Ingredient Name** 

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63779-575	
Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				

**Basis of Strength** 

Strength

Inactive Ingredients	
Ingredient Name	Strength
water (UNII: 059QF0KO0R)	
sodium laureth sulfate (UNII: BPV390UAP0)	
sodium lauryl sulfate (UNII: 368GB5141J)	
propylene glycol (UNII: 6DC9Q167V3)	
lauric diethanolamide (UNII: I29 I2VHG38)	
cocamidopropyl betaine (UNII: 5OCF3O11KX)	
coco diethanolamide (UNII: 92005F972D)	
citric acid monohydrate (UNII: 2968PHW8QP)	
dmdm hydantoin (UNII: BYR0546TOW)	
sodium chloride (UNII: 451W47IQ8X)	
FD&C Green NO. 3 (UNII: 3P3ONR6O1S)	

P	ackaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63779-575-05	3785 mL in 1 BOTTLE		
2	NDC:63779-575-42	50 mL in 1 BOTTLE		
3	NDC:63779-575-20	1100 mL in 1 BOTTLE		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	10/11/2010		

### Labeler - Canberra Corporation (068080621)

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

Revised: 8/2011 Canberra Corporation