

## **CARIBBEAN ESCAPE ANTI BACTERIAL FOAMING HAND - triclosan liquid**

### **Five Below**

*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 Fact**

#### **Active Ingredient**

Triclsan 0.3%

#### **Purpose**

Antiseptic

#### **Uses**

To decrease bacteria on the skin

#### **For external use only**

**When using this product: Keep out of eyes. Do not ingest.**

**Stop use and ask a doctor** if skin irritation develops

**Keep out of reach of children.** If swallowed, get medical assistance.

#### **Directions**

- Rub a dime size drop into hands

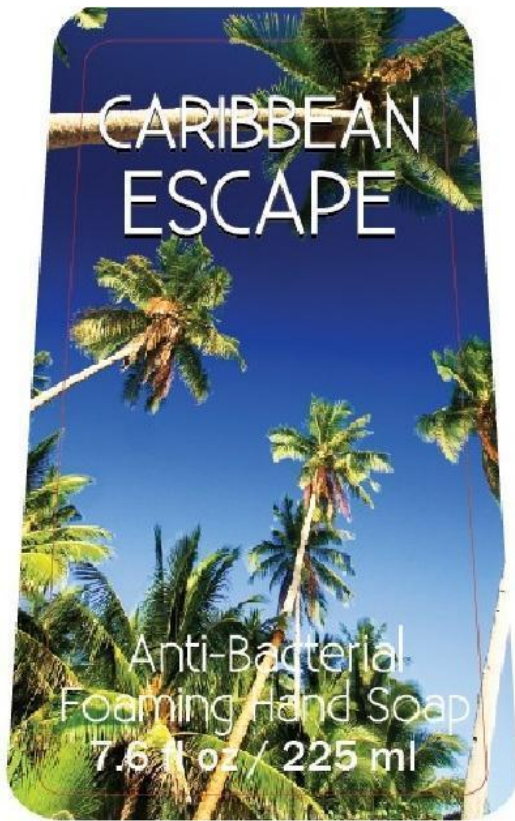
#### **Other Information:**

- Do not store above 105F
- May discolor some fabrics

#### **Inactive Ingredient:**

Deionized Water, Sodium Laurel Sulfate, Cocamide MEA, Fragrance, Citric Acid, Disodium EDTA, Methylisothiazolinone, Methylchloroisothiazolinone, may contain D and C Red No. 33, FD and C Red No. 40, FD and C Blue No. 1, FD and C Yellow No. 5, FD and C Yellow No. 6.

lab



Caribbean Escape

## CARIBBEAN ESCAPE ANTI BACTERIAL FOAMING HAND

triclosan liquid

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59021-002
Route of Administration	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TRICLOSAN (UNII: 4NM5039Y5X) (TRICLOSAN - UNII:4NM5039Y5X)	TRICLOSAN	0.3 g in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	82.009948 g in 100 mL
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
CO CO MONOETHANOLAMIDE (UNII: C80684146D)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	

ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59021-002-01	225 mL in 1 BOTTLE		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	04/13/2013	

**Labeler** - Five Below (095446659)

**Registrant** - Zhejiang Aidi Cosmetics Co., Ltd. (530876820)

### Establishment

Name	Address	ID/FEI	Business Operations
Zhejiang Aidi Cosmetics Co., Ltd.		530876820	manufacture(59021-002)

Revised: 4/2013

Five Below