SIMPLEPLEASURES SCENTED ANTI BACTERIAL HAND SANITIZER- alcohol liquid Tri-Coastal Design Company Inc.

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.

simplepleasures cranberry vanilla Hand Sanitizer

Active Ingredient

Alcohol 62%

Purpose

Antiseptic

Uses

- For handwashing to decrease bacteria on the skin
- Recommended for repeated use

Warnings

■ **For external use only** ■ Flammable, keep away from fire and flame ■ Does not contain grain alcohol; do not drink, if taken internally will produce serious gastric disturbances

When using this product

■ Avoid the eyes and mucous membranes ■ In the case of eyes or mucous membrane contact; rinse area thoroughly with water

Stop use and ask a doctor if

■ Condition worsens ■ Redness or irritation develops ■ If condition persists for more than 3 days

Keep out of reach of children

■ If swallowed, get medical help or Poison Control Center right away.

Directions

■ Rub dime sized amount between hands until dry ■ Supervise children in the use of this product ■ In the case of eye contact, rinse eyes thoroughly with water.

Other information

■ Store below 105°F ■ May discolor some fabrics

Inactive ingredients

Water, PEG-40 Hydrogenated Castor Oil, Fragrance, Acrylates/C10-30 Alkyl Acetate Crosspolymer,

Questions?

1-800-278-9218

Package Label





Drug Facts

(continued)

■ In the case of eyes or mucous membranes contact, rinse area thoroughly with water

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Drug Facts

(continued)

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Water, PEG-40 Hydrogenated Castor Oil, Fragrance, Acrylates (C10-30 Alkyl Acrylate Crosspolyme Sodium Hydroxide, FD&C Red No.40 (Cl 16035).

FD&C Red No.40 (Cl 16035), FD&C Blue No.1 (Cl 42090).

Questions?

■1-800-278-9218

Drug Facts

(continued)

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When using this product

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Drug Facts

(continued)

■ If swallowed contact a doctor or Poison Control Center immediately.

Directions

- Rub dime sized amount between hands until dry.
- Supervise children in the use of this product.
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alcohol liquid

Product Information	ict Information		
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49852-157
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 g in 100 mL

Inactive Ingredients	
Ingredient Name	Strength
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
WATER (UNII: 059QF0KO0R)	
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

P	nckaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49852-157- 01	30 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/12/2019	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/12/2019	

Labeler - Tri-Coastal Design Company Inc. (609734900)

Revised: 2/2020 Tri-Coastal Design Company Inc.