CRISP APPLE HAND SANI- ethyl alcohol liquid
Target 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.

Crisp Apple Hand Sanitizer
663.001/663AA/AB

Active ingredient

Ethyl Alcohol 65%

Purpose

Antiseptic

Use

- to decrease bcteria on the skin that could cause disease
- recommended for repeated use

Warnings

For external use only-hands

Flammable. Keep away from heat and flame.

When using this product

- keep out of eyes. Incase of contact with eyes, flush thoroughly with water.
- do not inhale or ingest
- avoid contact with broken skin

Stop use and ask a doctor

if skin irritation develops

Keep out of reach of children

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

Directions

- wet hands thoroughly with product and allow to dry without wiping
- for children under 6, use only under adult supervision

- not recommed for infants

Other Safety Information

- do not store above 105°F
- may discolor some fabrics
- harmful to wood finishes and plastics.

Inactive ingredients

benzophenone-4, carbomer, cellulose, fragrance, glycerin, hydroxypropyl methylcellulose, mannitol, red 4, red 40, retinyl palmitate, tocoperyl acetate, ultamarines, water

adverse reactions

Effective at eliminating 99.99% of many common harmful germs and bacteria in as little as 15 seconds

Made in U.S.A. with U.S. and foreign components

Dist. by Target Corp., Mpls., MN 55403

Questions or comments? 1-800-910-6874

Principal display panel

kills 99.99% of germs*

hand sanitizer

crisp apple

up & up

2 FL OZ (59.1 mL)

CRISP APPLE HAND SANI

ethyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-663
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	65 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
SULISOBENZONE (UNII: 1W6L629B4K)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
POWDERED CELLULOSE (UNII: SMD1X3XO9M)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MANNITOL (UNII: 3OWL53L36A)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ULTRAMARINE BLUE (UNII: I39WR998BI)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-663-16	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/05/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/05/2014	

Labeler - Target Corporation (006961700)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(11673-663)

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Vi-Jon, LLC		790752542	manufacture(11673-663)

Revised: 2/2023 Target Corporation