HAND SANITIZER- ethyl alcohol gel
Target Corp

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.

UP & UP Hand Sanitizer Kiwi Punch
544.000/544AA

Active ingredient

Ethyl alcohol 63%

Purpose

Antiseptic

Use

- to decrease bacteeria 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. In case 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

- irritation and redness develop
- condition persists for more than 72 hours

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 recommended for infants

Other information

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

Inactive ingredients

water, glycerin, tocopheryl acetate, retinyl palmitate, acrylates/C10-30 alkyl acrylate crosspolymer, benzophenone-4, mannitol, cellulose, hydroxypropyl methylcellulose, fragrance, yellow 5, blue 1, red 4, ultramarines

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

Questions or comments? 1-800-910-6874
Dist. by Target Corp., Mpls., MN
Made in U.S.A. with U.S. and foreign components
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PRINCIPAL DISPLAY PANEL

kills 99.99% of germs*

hand sanitizer kiwi punch

up & up

2 FL OZ (59.1 mL)

HAND SANITIZER
ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-086
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	567 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
glycerin (UNII: PDC6A3C0OX)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
SULISOBENZONE (UNII: 1W6L629B4K)	
MANNITOL (UNII: 3OWL53L36A)	
POWDERED CELLULOSE (UNII: SMD1X3XO9M)	
HYPROMELLOSE 2208 (100 MPA.S) (UNII: B1QE5P712K)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
ULTRAMARINE BLUE (UNII: I39WR998BI)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-086-16	59 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	03/14/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/14/2014	

Labeler - Target Corp (006961700)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(11673-086)

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Vi-Jon, LLC		790752542	manufacture(11673-086)

Revised: 3/2022 Target Corp