HAND SANITIZER- ethyl alcohol gel
Target Corp

UP & UP 544.000/544AA
Hand Sanitizer Kiwi Punch

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 Drug	505G(a)(3)	03/14/2014	

Labeler - Target Corp (006961700)

Registrant - Consumer Product Partners, LLC (119091520)

Establishment

Name	Address	ID/FEI	Business Operations
Consumer Product Partners, LLC		119091520	manufacture(11673-086)

Establishment

Name	Address	ID/FEI	Business Operations
Consumer Product Partners, LLC		119091514	manufacture(11673-086)