

**ADVANCED HAND SANITIZER- ethyl alcohol gel**  
**Target Corp**

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**Up & Up Advanced Hand Sanitizer with Aloe**  
**439.001/439AB-AC rev 1**

**Active ingredient**

Ethyl alcohol 70%

**Purpose**

Antiseptic

**Uses**

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

**Warnings**

**For external use only: hands**

**Flammable, keep away from heat or flame.**

**When using this product**

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

**Stop use and ask a doctor if**

- skin irritation develops
- 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

**Inactive ingredients**

water, aloe barbadensis leaf juice, glyceryl caprylate/caprate, glycerin, isopropyl myristate, tocopheryl acetate, acrylates/C10-30 alkyl acrylate crosspolymer, fragrance, benzophenone-4, blue 1, yellow 5

**Disclaimer**

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

Not tested on animals

**Adverse reaction**

049 00 0401 R00 C-002262-01-61-0149

Dist. by Target Corp. Mpls., MN 55403

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Guest Services 1-800-910-6874

**Principal display panel**

Hand Sanitizer

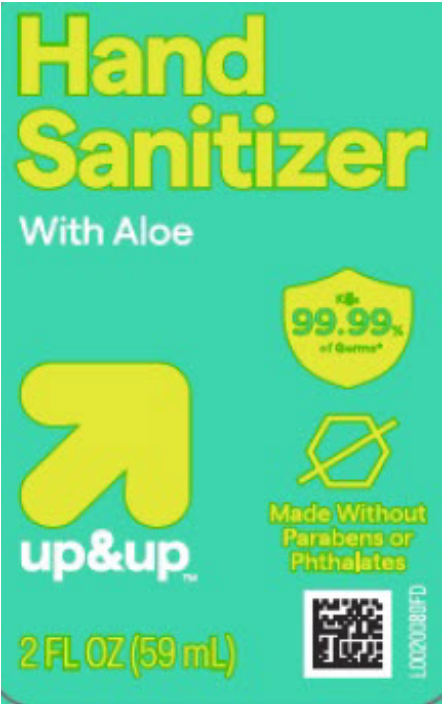
With Aloe

Kills 99.99% of germs\*

up + up <sup>TM</sup>

Made Without Parabens or Phthalates

2 FL OZ (59mL)



**ADVANCED HAND SANITIZER**

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-439
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)		ALCOHOL	70 mL in 100 mL
Inactive Ingredients			
Ingredient Name			Strength
WATER (UNII: 059QF0KO0R)			
ALOE VERA LEAF (UNII: ZY81Z83H0X)			
GLYCERYL CAPRYLATE/CAPRATE (UNII: G7515SW10N)			
GLYCERIN (UNII: PDC6A3C0OX)			
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)			
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)			
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)			
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)			
SULISOBENZONE (UNII: 1W6L629B4K)			
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)			
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-439-16	59 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	12/13/2012	
2	NDC:11673-439-34	236 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	12/13/2012	03/13/2022
3	NDC:11673-439-45	946 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	12/13/2012	02/24/2022
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug		505G(a)(3)	12/13/2012	

**Labeler** - Target Corp (006961700)

**Registrant** - Consumer Product Partners, LLC (119091520)

Establishment			
Name	Address	ID/FEI	Business Operations
Consumer Product Partners, LLC		119091520	manufacture(11673-439)

Establishment			
Name	Address	ID/FEI	Business Operations
Consumer Product Partners, LLC		119091514	manufacture(11673-439)