ADVANCED 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.

Advanced Hand Sanitizer with Aloe
439

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 and 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

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

*This product is not manufactured or distributed by GOJO Industries, Inc. distributor of Purell Refreshing Aloe Advanced Hand Sanitizer.

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

Questions? Call 1-800-910-6874
049-00-1701 ID 209446
Distributed by Target Corp.
Mpls., MN 55403
Made in U.S.A.

439.000/439AB

Principal Display Panel

advanced hand sanitizer gel with aloe

Compare to Purell**

More Effective Formula

Kills more than 99.99% of germs*

Hygiene handshake

up + up

2 FL OZ (59.1 mL)

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	700 mg in 1 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 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	
3	NDC:11673-439-45	946 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	12/13/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	12/13/2012	

Labeler - Target Corp (006961700)

Registrant - Vi-Jon (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon		088520668	manufacture(11673-439)

Revised: 5/2020 Target Corp