HAND SANITIZER WITH ALOE- 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.

Hand Sanitizer with Aloe

Active Ingredient

Alcohol 62%

Purpose

Antiseptic

Use

- Hand snitizer to help reduce bacteria that potentially can cause disease.
- For use when soap and water are not available.

Warnings

For external use only.

Flammable, Keep away from fire and flame

Do not use

- On childern less than 2 months of age.
- On open skin wounds.

When using this product

• Keep out of eyes, ears, and mouth. In case of conatct with eyes, rinse thoroughly with water.

Stop use and ask a doctor if irritation and redness develops and persists.

Keep out of reach of children

If swallowed, conatct a doctor or Poison Control Center right away.

Directions

- Apply a dime size amount to hands. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

Store between 59-86°F. Avoid freezing and excessive heat above 104°F.

Inactive ingredients

Water/Agua/Eau, Glycerin, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, T-Butyl Alcohol,

Aminomethyl Propanol, Aloe Barbadensis Leaf Extract, Tocopheryl Acetate, Denatonium Benzoate, FD&C Yellow No.5 (CI 19140), FD&C Blue No.1 (CI 42090).

Questions?

Call 1-800-910-6874

Distributed By

Target Corporation
Minneapolis, MN 55403

Package Label

Hand Sanitizer with Aloe

with silicone holder up & up 2 FL OZ (59.1 mL)





Drug Facts (continued)

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*Effective at eliminating more than 99.9% of many common bacteria in as little as

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HAND SANITIZER WITH ALOE

alcohol liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:11673-085

TOPICAL **Route of Administration**

Active Ingredient/Active Molety		
Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.62 mL in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)	
TERT-BUTYL ALCOHOL (UNII: MD83SFE959)	
AMINO METHYLPRO PANO L (UNII: LU49 E6626Q)	
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)	
.ALPHATO COPHERO L ACETATE (UNII: 9E8X80D2L0)	
DENATO NIUM BENZO ATE (UNII: 4YK5Z54AT2)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

I	Packaging				
#	t Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:11673-085- 16	1 in 1 CASE	12/18/2020		
1		59.1 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product			

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part333A	12/18/2020			

Labeler - Target Corporation (006961700)

Registrant - Fujian Haojin Toiletries Co Ltd. (554551930)

Establishment				
Name	Address	ID/FEI	Business Operations	
Fujian Haojin Toiletries Co Ltd.		554551930	manufacture(11673-085)	

Revised: 12/2020 Target Corporation