HAND SANITIZER- alcohol gel 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.

Trget Tricstl Gold Hnd Sntzr Bth Cncpts

Active Ingredient

Alcohol 62%

Purpose

Antiseptic

Use

■ For handwashing to decrease bacteria on the skin ■ Recommended for repeated use

Warnings

- For external use only
- Flammable, keep away from fire & flame
- Does not contain grain alcohol;do not drink. If taken internally will produce serious gastric disturbances

When using this product

- Avoid the eyes and mucous membranes
- In the case of eyes or mucous membranes contact, rinse area thoroughly with water

Stop use and ask a doctor if

- Condition worsens
- Redness or irritation develops
- If condition persists for more than 3 days

Keep out of reach of children

■ If swallowed contact a doctor or Poison Control Center immediately.

Directions

- Rub dime sized amount between hands until dry
- Supervise children in the use of this product
- In the case of eye contact, rinse eyes thoroughly with water

Other information

■ Store below 105°F ■ May discolor some fabrics

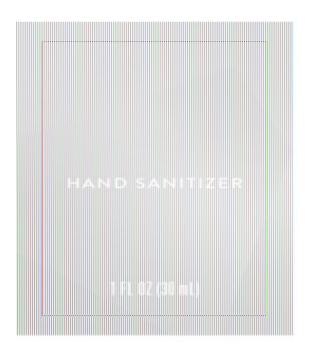
Inactive Ingredients

Water, Glycerin, Fragrance, Carbomer, Aminomethyl Propanol, Denatonium Benzoate, Mica, Yellow 5

Questions?

■ 1-800-278-9218

Package Label Gold





Drug Facts (continued)

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Drug Facts (continued)

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HAND SANITIZER

alcohol gel

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-256	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL	

Inactive Ingredients	
Ingredient Name	Strength
MICA (UNII: V8A1AW0880)	
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)	
WATER (UNII: 059QF0KO0R)	
DENATO NIUM BENZO ATE (UNII: 4YK5Z54AT2)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

ı	Packaging				
	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
	1 NDC:11673-256-	30 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/01/2020		

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part333A	07/01/2020			

Labeler - Target Corporation (006961700)

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