

DISNEY PRINCESS HAND SANITIZER- alcohol solution
Best Brand Consumers Products, Inc.

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.

Disney Princess Hand Sanitizer

Active Ingredient(s)

Ethyl Alcohol 68%. Purpose: Antiseptic

Purpose

Antiseptic

Use

To decrease the bacteria on the skin that could cause disease

Recommended for repeated use

Warnings

For external use only.

Flammable. Keep away from heat and flame

Discontinue if skin becomes irritated and ask a doctor

Keep out of eyes. In case of contact with eyes, flush thoroughly with water

Do not inhale or ingest

Avoid contact with broken skin

Keep out of reach of children. In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.

Directions

Wet hands thoroughly with products and rub until dry without wiping

For children under 6, use only under adult supervision

Not recommended for infants

Other information

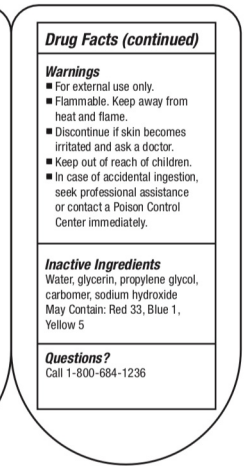
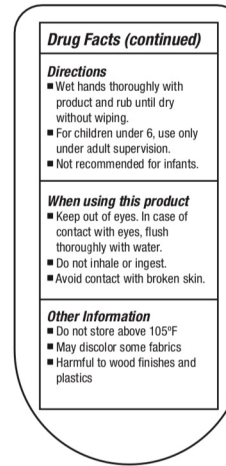
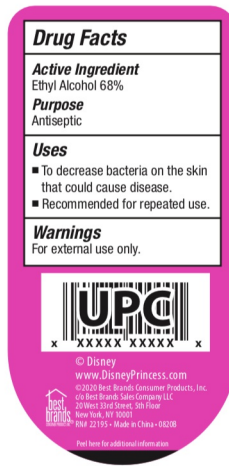
Do not store above 105F

May discolor some fabrics

Harmful to wood finishes and plastics

Inactive ingredients

Water, glycerin, propylene glycol, carbomer, sodium hydroxide. May contain: Red 33, Blue 1, Yellow 5



DISNEY PRINCESS HAND SANITIZER

alcohol solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74530-015
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	68 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
WATER (UNII: 059QF0K00R)	
CARBOMER 940 (UNII: 4Q93RCW27E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
FD&C BLUE NO. 1 (UNII: HBR47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74530-015-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/22/2020	
2	NDC:74530-015-02	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/22/2020	
3	NDC:74530-015-03	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/22/2020	
4	NDC:74530-015-04	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/22/2020	
5	NDC:74530-015-05	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/22/2020	

6	NDC:74530-015-06	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/22/2020	
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
OTC monograph not final	part333A		07/22/2020	

Labeler - Best Brand Consumers Products, Inc. (058304494)

Establishment			
Name	Address	ID/FEI	Business Operations
Ningbo Haishu Huayu Industry & Trade Co., Ltd.		527157032	manufacture(74530-015)

Revised: 7/2020

Best Brand Consumers Products, Inc.