# DISNEY FROZEN II 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.

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#### 74530-011

Drug Fact

# 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 assitance or contact a Poison Control Center immediately.

## **Directions**

Wet hands thoroughly with products and rub untill dry withour 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 wooden finishes and plastics

## **Inactive ingredients**

Water, glycerin, propylene glycol, carbomer, sodium hydroxide

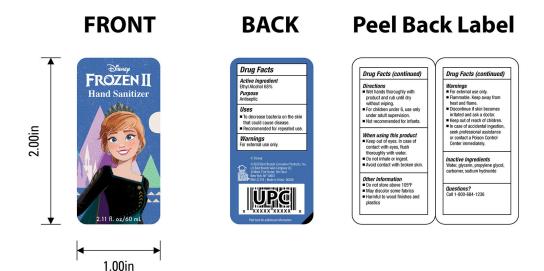
Package Label - Principal Display Panel

BEST BRANDS CONSUMER PRODUCTS Frozen 2

PRODUCT: Hand Sanitizer SKU: FRO2 HANS 01

Print Four Color Process





alcohol solution

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74530-011	
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)			
SO DIUM HYDRO XIDE (UNII: 55X0 4QC32I)			
WATER (UNII: 059QF0KO0R)			
CARBOMER 940 (UNII: 4Q93RCW27E)			
PROPYLENE GLYCOL (UNII: 6 DC9 Q16 7 V3)			

Packaging				
#	Item Code	Package Description	<b>Marketing Start Date</b>	Marketing End Date
1	NDC:74530-011-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/09/2020	
2	NDC:74530-011-02	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/09/2020	
3	NDC:74530-011-03	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/09/2020	
4	NDC:74530-011-04	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/09/2020	
5	NDC:74530-011-05	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/09/2020	
6	NDC:74530-011-06	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/09/2020	
7	NDC:74530-011-07	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/09/2020	
8	NDC:74530-011-08	90 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/09/2020	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	04/09/2020		

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

Establishment				
Name	Address	ID/FEI	<b>Business Operations</b>	
Ningbo Haishu Huayu Industry & Trade Co., Ltd.		527157032	manufacture (74530-011)	

Revised: 12/2020 Best Brand Consumers Products, Inc.