# HAND SANITIZER SAYFE- alcohol solution M3 Group LLC

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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#### Sayfe Hand Sanitizer

#### **Active Ingredient**

Alcohol 70% v/v

#### **Purpose**

Antiseptic

#### Use

■ hand sanitizer to help reduce bacteria on the skin. For use when soap and water are not available

#### **Warnings**

For external use only. Flammable. Keep away from fire or flame.

#### Do not use

■ in children less than 2 months of age ■ on open skin wounds

#### When using this product

■ avoid getting into the eyes. ■ in case of eye contact immediately flush eyes throughly with water

#### Stop use and ask a doctor if

■ irritation or redness develops ■ conditions persist for more than 72 hours ■ redness is present

#### Keep out of reach of children

In case of accidental ingestion, contact a doctor or Poison Control Center immediately.

#### **Directions**

■ place enough product in your palm to thoroughly cover both hands ■ rub hands together until dry ■ Supervise children under 6 years of age when using this product to avoid swallowing

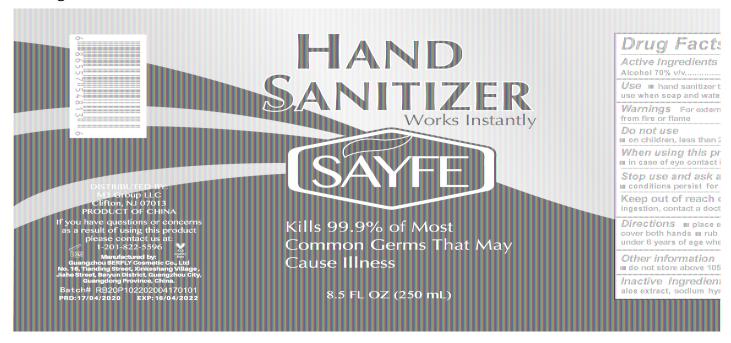
#### Other information

■ do not store above 105°F ■ may discolor fabrics or surfaces

#### **Inactive Ingredients**

water, glycerin, carbomer, aloe extract, sodium hydroxide

## **Package Label**



### HAND SANITIZER SAYFE

alcohol solution

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74176-001	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL		

Inactive Ingredients	
Ingredient Name	Strength
SO DIUM HYDRO XIDE (UNII: 55X0 4QC32I)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)	
WATER (UNII: 059QF0KO0R)	
CARBOMER HOMO POLYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)	

l	Packaging			
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:74176-001- 08	250 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/24/2020	
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Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	04/24/2020		

# Labeler - M3 Group LLC (104248779)

Establishment				
Name	Address	ID/FEI	Business Operations	
Guangzhou Berfly Cosmetic Co., Ltd		553098664	manufacture(74176-001)	

Revised: 4/2020 M3 Group LLC