KACHI HAND SANITIZER- alcohol gel
J&J SPIRITS

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.

Active ingredient

Alcohol 70% v/v

Purpose

Antiseptic

Use

Hand sanitizer 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 heat or flame.
Do not use
in children less than 2 months of age
on open skin wounds
When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.
Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients

glycerin, hydrogen peroxide, purified water USP

Product label

KACHI HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80219-001
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
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80219-001-01	30 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/21/2020	
2	NDC:80219-001-02	60 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/21/2020	
3	NDC:80219-001-03	120 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/21/2020	
4	NDC:80219-001-04	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/21/2020	
5	NDC:80219-001-05	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/21/2020	
6	NDC:80219-001-06	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/21/2020	
7	NDC:80219-001-07	1892 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/21/2020	
8	NDC:80219-001-08	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/21/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/21/2020	

Labeler - J&J SPIRITS (815978244)

Revised: 8/2020 J&J SPIRITS