HAND SANITIZER- ethyl alcohol gel
A. J. & W. Incorporated

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(s)

Ethyl Alcohol 75%

Purpose

Antiseptic, Hand Sanitizer

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.
- Orally.
- For other purposes other than specified.

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.

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Directions

Wet hands thoroughly and 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

Water, Glycerin, Carbomer, Aloe Vera Leaf

Package Label - Principal Display Panel

100 ml NDC: 74839-001-10

500 ml NDC: 74839-001-50

1000 ml NDC: 74839-001-00

HAND SANITIZER

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:74839-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75.59 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOMER 980 (UNII: 4Q93RCW27E)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:74839-001-10	100 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/14/2020	
2	NDC:74839-001-50	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/14/2020	
3	NDC:74839-001-00	1000 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/14/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/14/2020	

Labeler - A. J. & W. Incorporated (064157654)

Registrant - A. J. & W. Incorporated (064157654)

Revised: 5/2020 A. J. & W. Incorporated