ACP ANTIBACTERIAL ALCOHOL WIPES- ethyl alcohol cloth Swanrose, 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.

ACP Antibacterial Alcohol Wipes

Drug Facts

Active Ingredient

Ethyl Alcohol 70%

Purpose

Antiseptic Handwash

Uses

· For hand washing to decrease bacteria on the skin.

Store in a cool, dry place

Directions

· Wet hands thoroughly with product and allow to dry without wiping.

Other Information

- · Do not store above 110°F
- · May discolor some fabrics

Warnings

For external use only,

Do not use- in mouth or touch eyes

Non-flus hable

Flammable - Keep away from fire or flame

When using this product, avoid contact with eyes. In case of eye contact, rinse with water to remove.

Stop use and ask a doctor if irritation and redness develop.

Keep out of reach of children

Inactive Ingredients

Aqua, Glycerin, Aloe Barbadensis Leaf Juice.

With 70% Alcohol

- $\sqrt{\mathbf{Essential}}$ for home or outside
- $\sqrt{\text{Contains alcohol}}$
- $\sqrt{\text{Used for hands and surface}}$

Marketed & distributed by Swanrose Inc.

Address:SWANROSE, INC.

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LOS ANGELES, CA 90046

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Shelf Life: 24 months





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- ✓ Used for hands and surface

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ACP ANTIBACTERIAL ALCOHOL WIPES

ethyl alcohol cloth

Product Information

Product TypeHUMAN OTC DRUGItem Code (Source)NDC:78287-140Route of AdministrationTOPICAL

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)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:78287-140- 02	1 in 1 POUCH	06/25/2020		
1		3.57 mL in 1 PACKAGE; Type 0: Not a Combination Product			
2	NDC:78287-140- 03	10 in 1 POUCH	06/25/2020		
2		3.57 mL in 1 PACKAGE; Type 0: Not a Combination Product			
3	NDC:78287-140-01	50 in 1 POUCH	06/25/2020		
3		3.57 mL in 1 PACKAGE; Type 0: Not a Combination Product			

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not final	part333A	06/25/2020				

Labeler - Swanrose, Inc. (117523391)

Revised: 7/2020 Swanrose, Inc.