PURELL ALCOHOL FORMULATION- alcohol swab
GOJO Industries, 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.

PURELL Hand Sanitizing Wipes Alcohol Formulation

Active ingredient

Ethyl alcohol 62%

Purpose

Antimicrobial

Use

Hand sanitizer to help reduce bacteria on the skin

Warnings

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

When using this product do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash appears and lasts

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

• Wet hands thoroughly with product and allow to dry

• Children under 6 years of age should be supervised when using PURELL® products

Other information

• Store below 110°F (43°C)

• May discolor certain fabrics or surfaces

Inactive ingredients

Water (Aqua), Isopropyl Alcohol, Glycerin, Aloe Barbadensis Leaf Juice, Isopropyl Myristate, Propylene Glycol, Retinyl Palmitate, Tocopheryl Acetate, Zea Mays (Corn) Oil

PURELL ALCOHOL FORMULATION

alcohol swab

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:21749-367
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.62 mL in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
CORN OIL (UNII: 8470G57WFM)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:21749-367-76	4 mL in 1 POUCH; Type 0: Not a Combination Product	02/24/2012	
2	NDC:21749-367-24	96 mL in 1 PACKAGE; Type 0: Not a Combination Product	02/24/2012	
3	NDC:21749-367-36	144 mL in 1 PACKAGE; Type 0: Not a Combination Product	02/24/2012	
4	NDC:21749-367-63	315 mL in 1 CANISTER; Type 0: Not a Combination Product	02/24/2012	
5	NDC:21749-367-94	540 mL in 1 PACKAGE; Type 0: Not a Combination Product	02/24/2012	
6	NDC:21749-367-68	700 mL in 1 CANISTER; Type 0: Not a Combination Product	02/24/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	02/24/2012	

Labeler - GOJO Industries, Inc. (004162038)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	manufacture(21749-367)

Revised: 12/2017 GOJO Industries, Inc.