

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



**TOUGH ON MESSES,
GENTLE ON HANDS**

***Kills 99.99% of most common germs that may make you sick**

- Helps reduce the risk of infections
- Leaves no stickiness or residue on hands
- With moisturizers and vitamins A & E
- Removes light soils and dirt from hands
- An antiseptic hand sanitizer to help prevent cross-contamination
- Hypoallergenic. Dermatologist tested; non-irritating
- Paraben-free, fragrance-free, dye-free formulation
- Recommended for repeated use
- Food Code Compliant — Meets Food Code Hand Sanitizers criteria (Section 2-301.16) published by the FDA

Convenient for use in healthcare, nursing stations, patient bedside, long-term nursing care, the home, office, car, restaurants, health clubs, workshops ... whenever soap and water are not available.



Distributed by:
GOJO Industries, Inc.
Akron, OH 44309
Questions?
Tel: 1-888-4-PURELL
www.PURELL.com
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9030-643

To Open Package:

- Flip open dispensing cap and remove entire lid from container by lifting upward.
- Locate wipe at center of roll, twist corner to a point and thread through small opening in lid. DO NOT PUSH FINGER THROUGH OPENING.



- Replace lid and pull sheet up at an angle. Remaining wipes feed automatically.
- When finished, snap lid cap shut to retain moisture.

Drug Facts

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Questions?
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• www.GOJO.com



HAND SANITIZING WIPES

**TOUGH ON MESSES,
GENTLE ON HANDS**



**KILLS 99.99%
OF ILLNESS CAUSING GERMS***

DERMATOLOGIST TESTED • PARABEN FREE

**FRAGRANCE FREE
ALCOHOL FORMULA**

**80
WIPES**

Wipe Dimensions:
6 in. x 7 in.
(15.2 cm x 17.8 cm)

Reorder No. 9030-12



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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)