PURELL HAND SANITIZING WIPE- alcohol cloth
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 Wipe

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

Wipe hands; discard

Inactive ingredients

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

Purell®

HAND SANITIZING

Wipe

KILLS 99.99%
OF MOST ILLNESS CAUSING GERMS*

FRAGRANCE FREE

ONE WIPE • ALCOHOL FORMULA

Drug Facts X

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Other information May discolor certain fabrics or surfaces

Paraben free • Phthalate free

***Kills 99.99% of most common germs that may cause illness**

Distributed by: GOJO Industries, Inc. Akron OH 44309

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PURELL HAND SANITIZING WIPE

alcohol cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:21749-363
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
GLYCERIN (UNII: PDC6A3C0OX)	
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:21749-363-76	1 in 1 PACKAGE	01/22/2007	
1		1.97 mL in 1 POUCH; 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	part333E	01/22/2007	

Labeler - GOJO Industries, Inc. (004162038)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	manufacture(21749-363)