PURELL ADVANCED REFRESHING- alcohol gel
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 Advanced Hand Sanitizer Refreshing Gel

Active ingredient

Ethyl alcohol 70% v/v

Purpose

Antimicrobial

Uses

- 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

- Put enough product in your palm to cover hands and rub hands together briskly until dry
- Children under 6 years of age should be supervised when using PURELL

Other information

- Store below 110°F (43°C)
- May discolor certain fabrics or surfaces

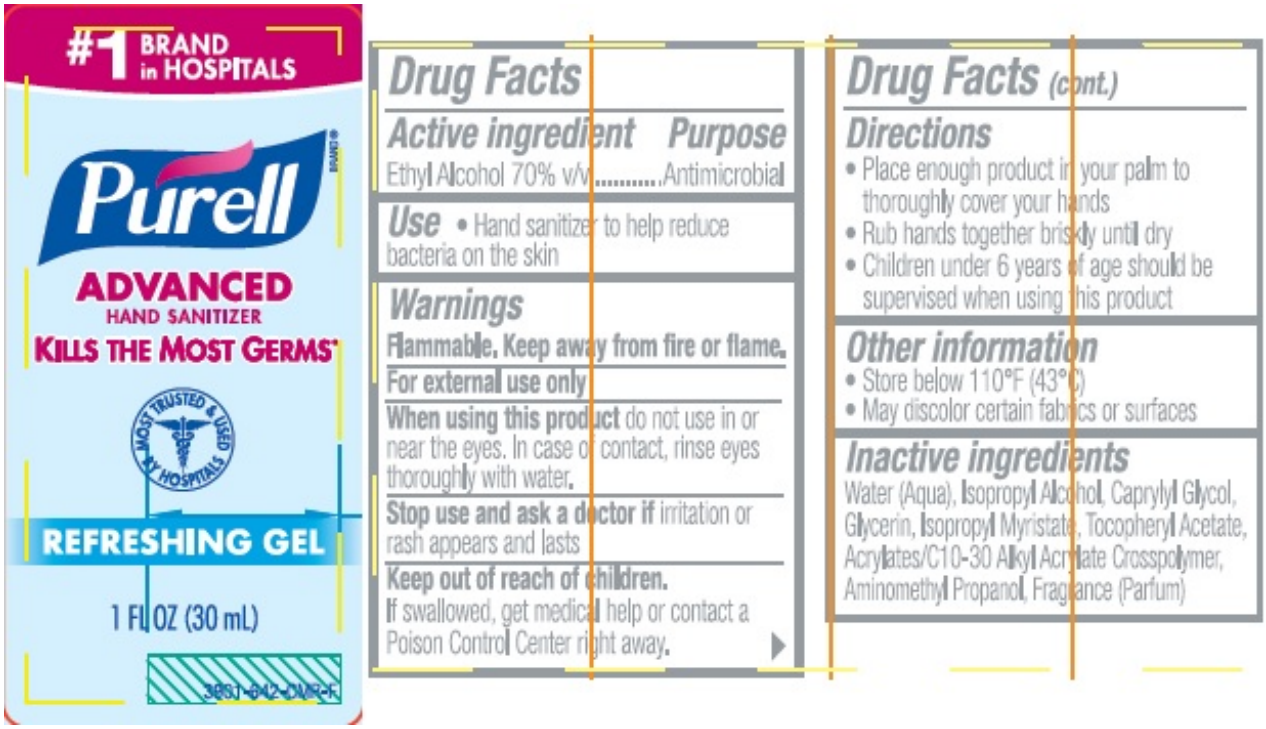
Inactive ingredients

Water (Aqua), Isopropyl Alcohol, Caprylyl Glycol, Glycerin, Isopropyl Myristate, Tocopheryl Acetate, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aminomethyl Propanol, Fragrance (Parfum)

Distributed by:
GOJO Industries, Inc.

Akron, OH 44309
Questions?
Tel: 1-888-4-PURELL
www.PURELL.com

Made in U.S.A.



PURELL ADVANCED REFRESHING

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:21749-704
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.7 mL in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	

.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21749-704-01	29 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/15/2012	
2	NDC:21749-704-02	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/15/2012	
3	NDC:21749-704-08	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/15/2012	
4	NDC:21749-704-12	354 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/15/2012	
5	NDC:21749-704-04	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/15/2012	
6	NDC:21749-704-10	295 mL in 1 PACKAGE; Type 0: Not a Combination Product	03/15/2012	
7	NDC:21749-704-20	2000 mL in 1 PACKAGE; Type 0: Not a Combination Product	03/15/2012	
8	NDC:21749-704-59	591 mL in 1 PACKAGE; Type 0: Not a Combination Product	03/15/2012	
9	NDC:21749-704-50	15 mL in 1 PACKAGE; Type 0: Not a Combination Product	03/15/2012	
10	NDC:21749-704-45	450 mL in 1 PACKAGE; Type 0: Not a Combination Product	03/15/2012	
11	NDC:21749-704-33	1000 mL in 1 PACKAGE; Type 0: Not a Combination Product	03/15/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/15/2012	

Labeler - GOJO Industries, Inc. (004162038)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	MANUFACTURE(21749-704)

Revised: 1/2017 GOJO Industries, Inc.