PURELL ADVANCED REFRESHING ALOE- 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 Aloe

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.

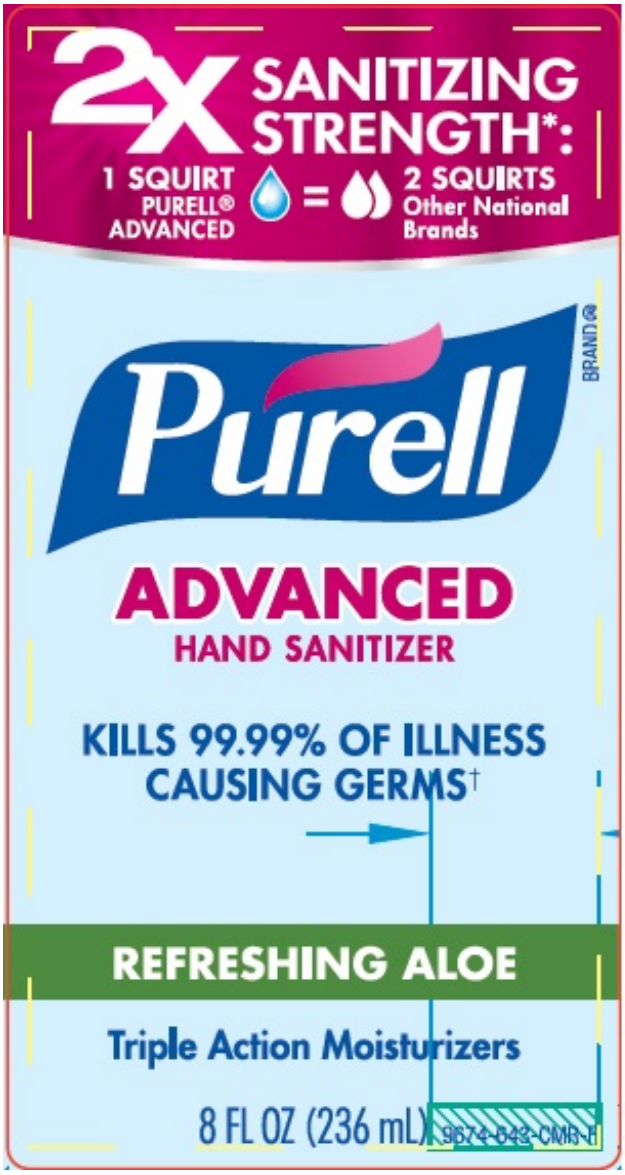
Directions

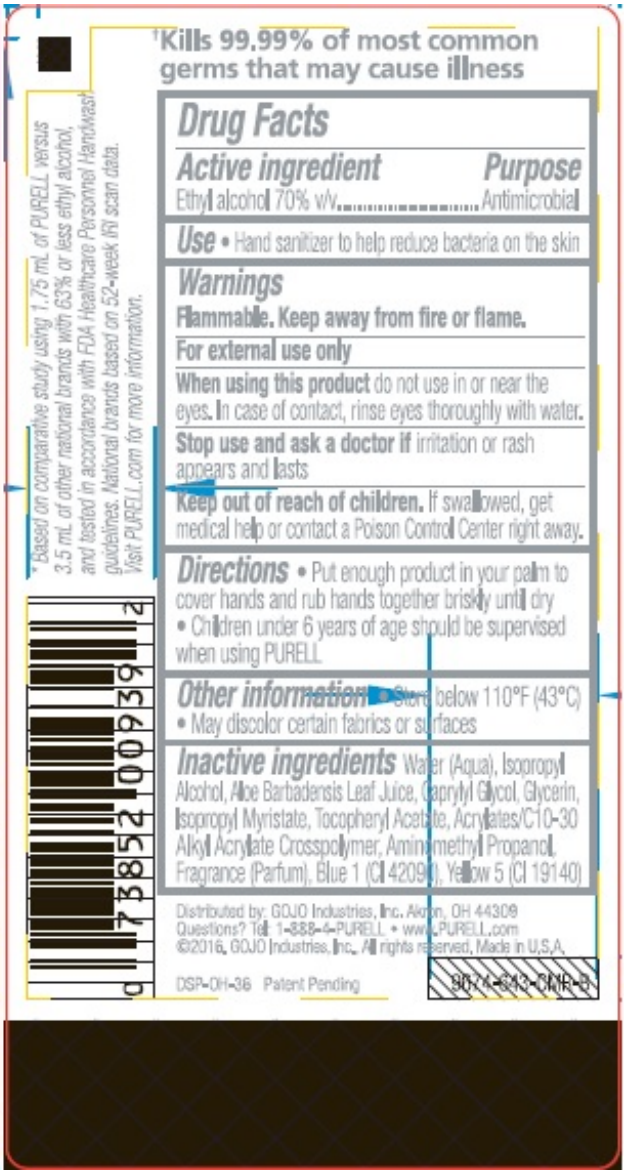
- Put enough product in your palm to cover hands and rub hands together briskly until dry
- Children under 6 years of age should be supervised when using PURELL

Inactive ingredients

Water (Aqua), Isopropyl Alcohol, Aloe Barbadensis Leaf Juice, Caprylyl Glycol, Glycerin, Isopropyl Myristate, Tocopheryl Acetate, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aminomethyl Propanol, Fragrance (Parfum), Blue 1 (CI 42090), Yellow 5 (CI 19140)

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


2X SANITIZING STRENGTH*:
1 SQUIRT PURELL® ADVANCED = 2 SQUIRTS Other National Brands
Purell
BRAND
ADVANCED
HAND SANITIZER
KILLS 99.99% OF ILLNESS CAUSING GERMS†
REFRESHING ALOE
Triple Action Moisturizers
8 FL OZ (236 mL)
9674-843-CMR-H


†Kills 99.99% of most common germs that may cause illness
* based on comparative study using 1.75 mL of PURELL versus 3.5 mL of other national brands with 63% or less ethyl alcohol, and tested in accordance with FDA Healthcare Personnel Handwash guidelines. National brands based on 52-week IRI scan data. Visit PURELL.com for more information.
Drug Facts
Active ingredient Purpose
Ethyl alcohol 70% v/v ..... 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 • 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, Aloe Barbadensis Leaf Juice, Caprylyl Glycol, Glycerin, Isopropyl Myristate, Tocopheryl Acetate, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aminomethyl Propanol, Fragrance (Parfum), Blue 1 (CI 42090), Yellow 5 (CI 19140)
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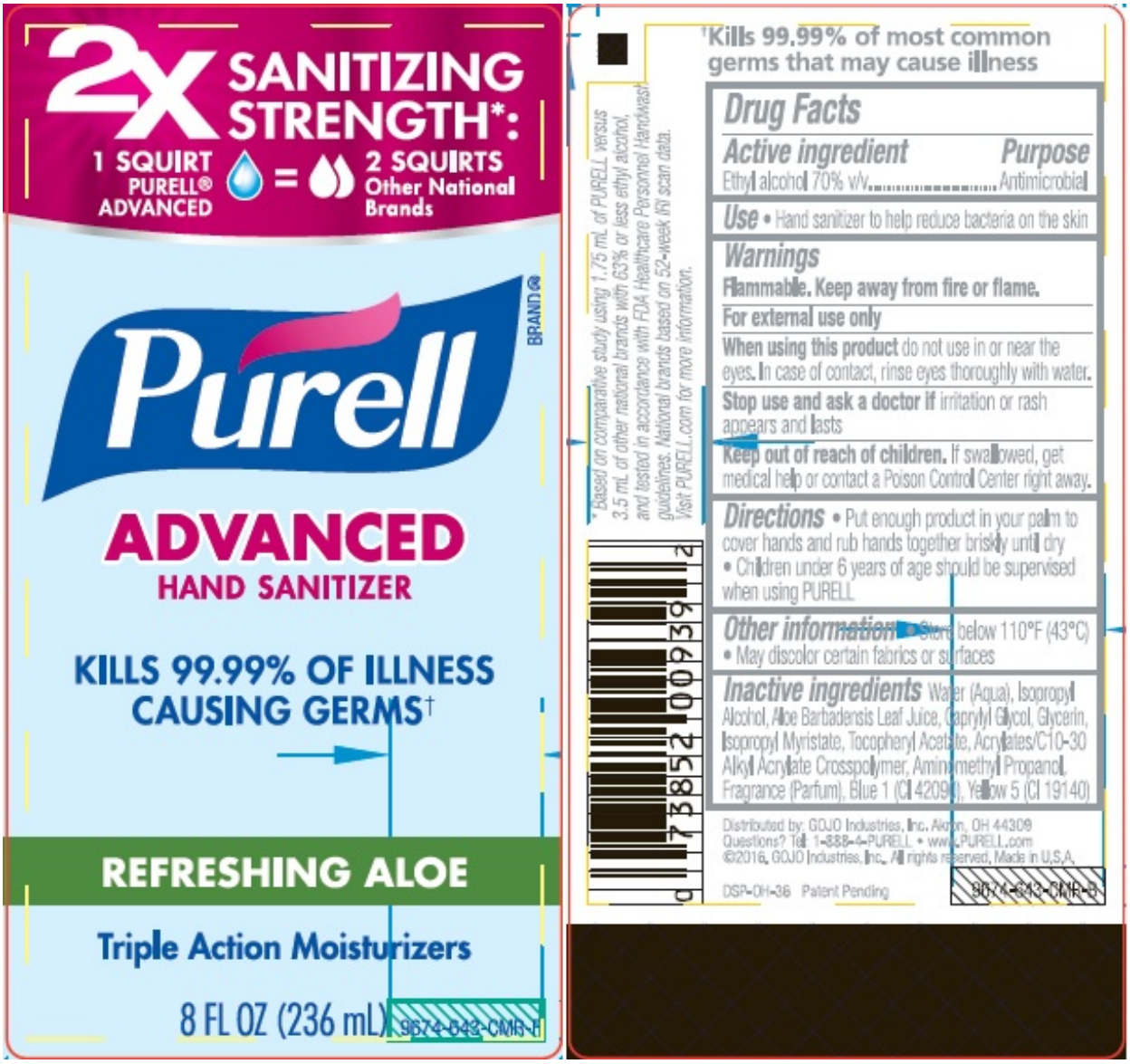
DSP-OH-36 Patent Pending
9674-843-CMR-H



PURELL ADVANCED REFRESHING ALOE

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:21749-705
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)	

ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21749-705-50	15 mL in 1 PACKAGE; Type 0: Not a Combination Product	03/14/2012	01/01/2022
2	NDC:21749-705-01	30 mL in 1 PACKAGE; Type 0: Not a Combination Product	03/14/2012	01/01/2022
3	NDC:21749-705-02	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/14/2012	01/01/2022
4	NDC:21749-705-04	118 mL in 1 PACKAGE; Type 0: Not a Combination Product	03/14/2012	01/01/2022
5	NDC:21749-705-08	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/14/2012	01/01/2022
6	NDC:21749-705-10	295 mL in 1 PACKAGE; Type 0: Not a Combination Product	03/14/2012	01/01/2022
7	NDC:21749-705-12	354 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/14/2012	01/01/2022
8	NDC:21749-705-20	591 mL in 1 PACKAGE; Type 1: Convenience Kit of Co-Package	03/14/2012	01/01/2022
9	NDC:21749-705-80	800 mL in 1 PACKAGE; Type 0: Not a Combination Product	03/14/2012	01/01/2022
10	NDC:21749-705-33	1000 mL in 1 PACKAGE; Type 0: Not a Combination Product	03/14/2012	01/01/2022
11	NDC:21749-705-89	1200 mL in 1 PACKAGE; Type 0: Not a Combination Product	03/14/2012	01/01/2022

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/14/2012	01/01/2022

Labeler - GOJO Industries, Inc. (004162038)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	MANUFACTURE(21749-705)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		088312414	MANUFACTURE(21749-705) , label(21749-705) , pack(21749-705)