

PURELL ADVANCED HAND SANITIZER BE VIBRANT- 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 be Vibrant

Active ingredient

Ethyl alcohol 70% v/v

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

- Place enough product in your palm to thoroughly cover your hands
- Rub hands together briskly until dry
- Children under 6 years of age should be supervised when using this product

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), Blue 1 (CI 42090), Yellow 5 (CI 19140)



Drug Facts

Active ingredient	Purpose
Ethyl Alcohol 70% w/v	Antimicrobial

Use • Hand sanitizer to help reduce bacteria on the skin

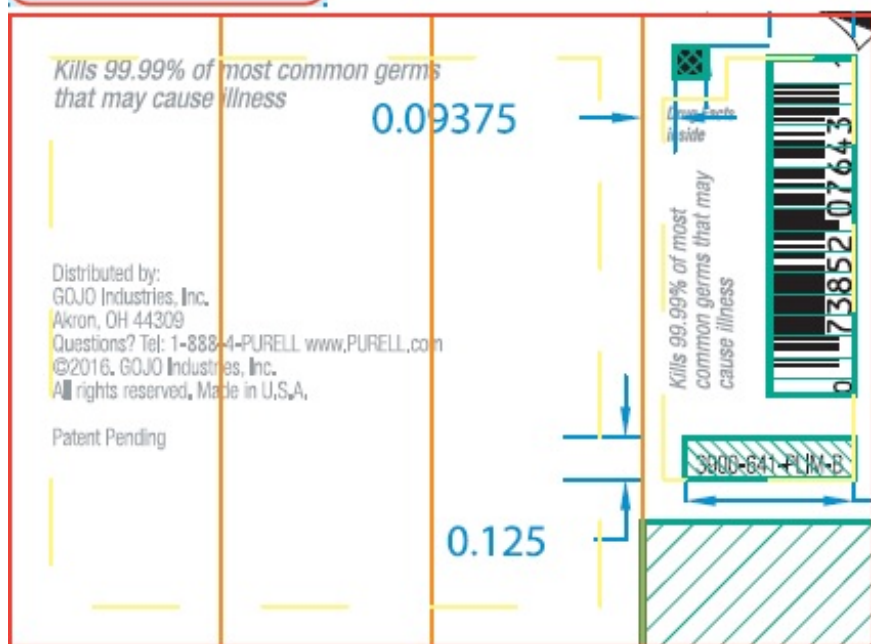
Warnings
Flammable. Keep away from fire or flame.
For external use only.
When using this product do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.
Stop use and ask a doctor if irritation or rash appears and lasts
Keep out of reach of children.
If swallowed, get medical help or contact a Poison Control Center right away.

Drug Facts (continued)

Directions • Place enough product in your palm to thoroughly cover your hands
 • Rub hands together briskly until dry
 • Children under 6 years of age should be supervised when using this product

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), Blue 1 (CI 42090), Yellow 5 (CI 19140)



PURELL ADVANCED HAND SANITIZER BE VIBRANT

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:21749-028
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 (UNII: 9E8X80D2L0)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
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:21749-028-01	30 mL in 1 PACKAGE; Type 0: Not a Combination Product	12/22/2016	
2	NDC:21749-028-02	59 mL in 1 PACKAGE; Type 0: Not a Combination Product	12/22/2016	
3	NDC:21749-028-04	118 mL in 1 PACKAGE; Type 0: Not a Combination Product	12/22/2016	
4	NDC:21749-028-08	236 mL in 1 PACKAGE; Type 0: Not a Combination Product	12/22/2016	
5	NDC:21749-028-10	295 mL in 1 PACKAGE; Type 0: Not a Combination Product	12/22/2016	
6	NDC:21749-028-12	354 mL in 1 PACKAGE; Type 0: Not a Combination Product	12/22/2016	
7	NDC:21749-028-45	450 mL in 1 PACKAGE; Type 0: Not a Combination Product	12/22/2016	
8	NDC:21749-028-16	473 mL in 1 PACKAGE; Type 0: Not a Combination Product	12/22/2016	
9	NDC:21749-028-20	591 mL in 1 PACKAGE; Type 0: Not a Combination Product	12/22/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	12/22/2016	

Labeler - GOJO Industries, Inc. (004162038)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	manufacture(21749-028)

Revised: 12/2016

GOJO Industries, Inc.