PURELL ADVANCED HAND SANITIZER BE VIBRANT- alcohol gel
GOJO Industries, Inc.

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 shodl be supervised when using this product

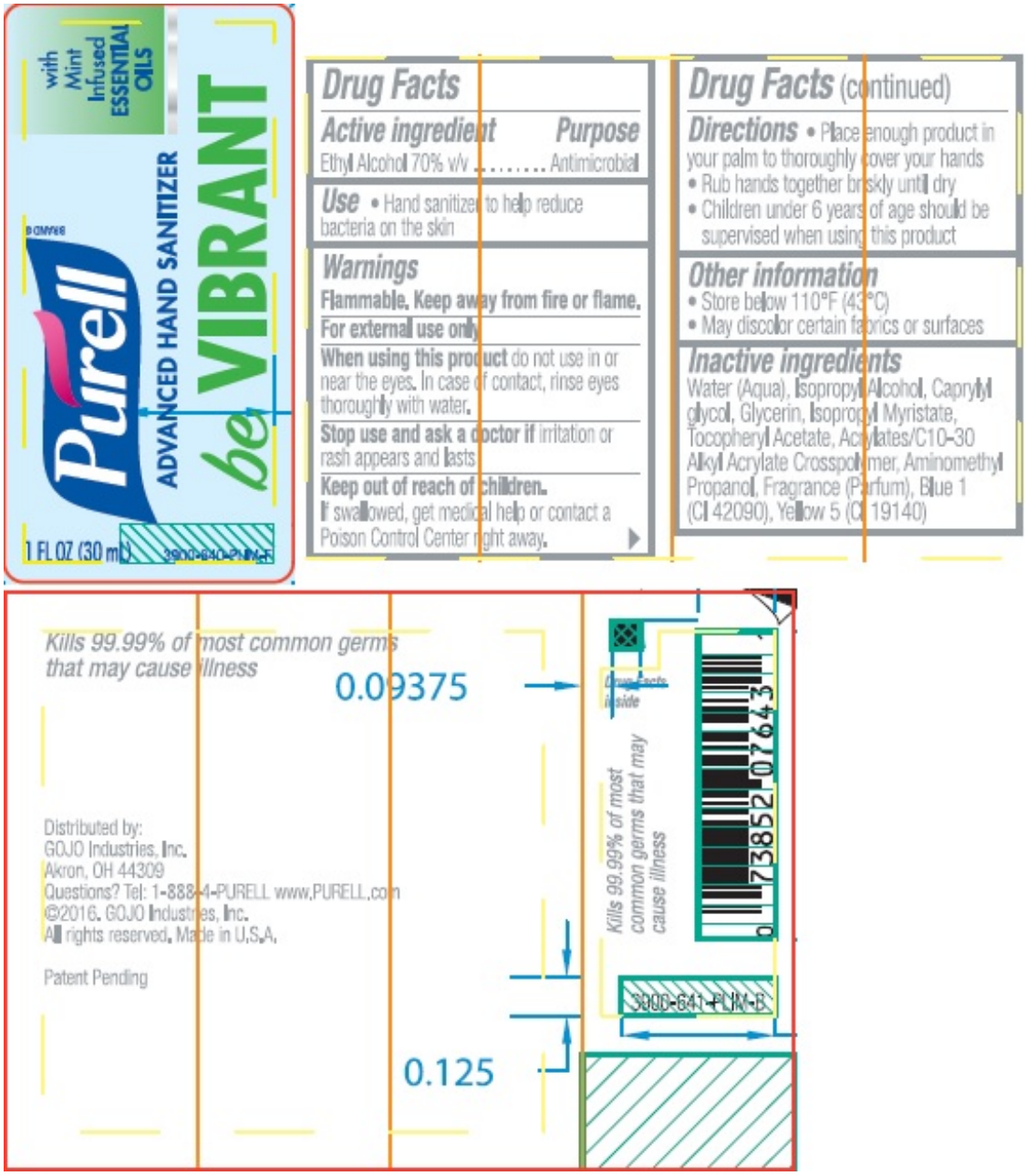
Other information

• 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	12/31/2024
3	NDC:21749-028-04	118 mL in 1 PACKAGE; Type 0: Not a Combination Product	12/22/2016	12/31/2024
4	NDC:21749-028-08	236 mL in 1 PACKAGE; Type 0: Not a Combination Product	12/22/2016	12/31/2024
5	NDC:21749-028-10	295 mL in 1 PACKAGE; Type 0: Not a Combination Product	12/22/2016	12/31/2024
6	NDC:21749-028-12	354 mL in 1 PACKAGE; Type 0: Not a Combination Product	12/22/2016	12/31/2024
7	NDC:21749-028-45	450 mL in 1 PACKAGE; Type 0: Not a Combination Product	12/22/2016	12/31/2024
8	NDC:21749-028-16	473 mL in 1 PACKAGE; Type 0: Not a Combination Product	12/22/2016	12/31/2024
9	NDC:21749-028-20	591 mL in 1 PACKAGE; Type 0: Not a Combination Product	12/22/2016	12/31/2024

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	12/22/2016	

Labeler - GOJO Industries, Inc. (004162038)

Establishment

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

Revised: 1/2026 GOJO Industries, Inc.