

**PURELL ADVANCED HAND SANITIZER NATURALS GEL- alcohol gel**  
**GOJO Industries, Inc.**

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**PURELL Advanced Hand Sanitizer Naturals Gel**

**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**

- 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, Caprylyl glycol, Citrus Aurantium Dulcis (Orange) Peel Oil, Glycerin, Isopropyl Myristate, Lavandula Hybrida (Lavandin) oil, Litsea Cubeba Fruit Oil, Pelargonium Graevolens (Geranium) oil, Pogostemon Cablin Oil, Tocopheryl Acetate, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aminomethyl Propanol



**ADVANCED  
HAND SANITIZER**



**NATURALS**  
MADE WITH PLANT BASED ALCOHOL

**Kills 99.99% of Germs\***  
With Essential Oils  
and Skin Conditioners

6.5 FL OZ (192 mL) 3265-640-CMR-F



**ADVANCED  
HAND SANITIZER**

**NATURALS**

100% naturally renewable ethanol supports well-being for your family and your world.

- Ingredients are 93% naturally derived
- Naturally fragrances
- Triclosan, paraben, and preservative free

**Active ingredient**  
Ethyl Alcohol 70% v/v

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

Distributed by: 6030 Industries, Inc., Akron, OH 44308  
Questions? Tel: 1-888-4-PURELL www.PURELL.com  
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Made in the USA with U.S. and foreign components  
3265-640-CMR-F

0 73852 29352 4

**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.

HINGE  
HINGE

**Drug Facts (cont.)**

**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, Citrus Aurantium Dulcis (Orange) Peel Oil, Glycerin, Isopropyl Myristate, Lavandula Hybrid (Lavandin) oil, Litsea Cubeba Fruit Oil, Pelargonium Graevolens (Geranium) oil, Pogostemon Cablin Oil, Tocopheryl Acetate, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aminomethyl Propanol

alcohol gel

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:21749-625
<b>Route of Administration</b>	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)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
BITTER ORANGE OIL (UNII: 9TLV70SV6I)	
LAVANDIN OIL (UNII: 9RES347CKG)	
LITSEA OIL (UNII: 2XIW34BN6O)	
GERANIUM OIL, ALGERIAN TYPE (UNII: 5Q1I94P4WG)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
PATCHOULI OIL (UNII: F3IN55X5PO)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21749-625-01	30 mL in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2016	
2	NDC:21749-625-02	59 mL in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2016	
3	NDC:21749-625-04	118 mL in 1 PACKAGE; Type 0: Not a Combination Product	12/15/2019	12/31/2023
4	NDC:21749-625-08	236 mL in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2016	
5	NDC:21749-625-29	295 mL in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2016	
6	NDC:21749-625-12	355 mL in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2016	
7	NDC:21749-625-28	828 mL in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2016	
8	NDC:21749-625-10	1000 mL in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2016	
9	NDC:21749-625-24	710 mL in 1 PACKAGE; Type 0: Not a Combination Product	08/01/2021	
10	NDC:21749-	192 mL in 1 PACKAGE; Type 0: Not a Combination	07/01/2021	

10	625-65	Product	07/01/2021
<b>Marketing Information</b>			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	01/01/2016	

**Labeler -** GOJO Industries, Inc. (004162038)

<b>Establishment</b>			
Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	manufacture(21749-625)

<b>Establishment</b>			
Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		088312414	manufacture(21749-625) , label(21749-625) , pack(21749-625)

Revised: 12/2025

GOJO Industries, Inc.