

**PURELL ADVANCED E3 RATED INSTANT HAND SANITIZER- alcohol liquid**  
**GOJO Industries, Inc.**

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**PURELL Advanced E3 Rated Instant Hand Sanitizer**

**Active ingredient**

Ethyl Alcohol 70% v/v

**Purpose**

Antimicrobial

**Uses**

- Hand sanitizer to help reduce bacteria on the skin that could cause disease
- Recommended for repeated use

**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
- No rinsing required
- No towels needed

**Inactive ingredients**

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

**Drug Facts**

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**Other information**

- Do not store above 110°F (43°C) ■ May discolor certain fabrics or surfaces

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**Questions or comments?**

Call 1-800-321-9647 Monday through Friday 8:00 AM to 5:00 PM



ADVANCED  
**E3 Rated**  
INSTANT **HAND**  
**SANITIZER**

FOR USE IN FOOD PROCESSING

AVANZADOS  
**Clasificación E3**  
ANTISÉPTICO **INSTANTÁNEO**  
**PARA LAS MANOS**  
PARA EL USO EN PROCESAMIENTO  
DE ALIMENTO

Distributed by: Distribuido por:  
**GOJO Industries, Inc.** Akron, OH 44309  
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Tel: 800-321-9647 • 330-255-6000  
www.GOJO.com  
Made in U.S.A., Hecho en los E.E.U.U.

Refill for the PURELL® NXT® SPACE SAVER™  
System • Sistema  
Patent Pending  
Patente Pendiente



U.S. Pat. 6,216,916  
U.S. Pat. D432,547  
U.S. Pat. 6,619,512  
U.S. Pat. 6,877,642  
U.S. Pat. D431,404

2163

1 L 33.8 FL OZ



Nonfood Compounds  
Program Listed E3 145336

**PURELL ADVANCED E3 RATED INSTANT HAND SANITIZER**

alcohol liquid

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:21749-706
<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

<b>WATER</b> (UNII: 059QF0KO0R)	
<b>ISOPROPYL ALCOHOL</b> (UNII: ND2M416302)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>ISOPROPYL MYRISTATE</b> (UNII: 0RE8K4LNJS)	
<b>.ALPHA.-TOCOPHEROL ACETATE, D-</b> (UNII: A7E6112E4N)	
<b>CAPRYLYL GLYCOL</b> (UNII: 00YIU5438U)	
<b>AMINOMETHYLPROPANOL</b> (UNII: LU49E6626Q)	

<b>Packaging</b>				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21749-706-08	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/31/2013	12/31/2024
2	NDC:21749-706-12	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/31/2013	12/31/2024
3	NDC:21749-706-97	700 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/31/2013	12/31/2024
4	NDC:21749-706-10	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/31/2013	12/31/2024
5	NDC:21749-706-89	1200 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/31/2013	

<b>Marketing Information</b>			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	03/31/2013	

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

<b>Establishment</b>			
Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	manufacture(21749-706)

<b>Establishment</b>			
Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		088312414	manufacture(21749-706) , label(21749-706) , pack(21749-706)