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

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

PURELL ADVANCED E3 RATED INSTANT HAND SANITIZER

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:21749-706
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)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	

Packaging

#	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	

Marketing Information

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)

Establishment

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

Establishment

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

Revised: 1/2026 GOJO Industries, Inc.