

**PURELL ADVANCED HAND SANITIZER ULTRA NOURISHING FOAM-
alcohol liquid
GOJO Industries, Inc.**

PURELL Advanced Hand Sanitizer ULTRA NOURISHING Foam

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

Inactive ingredients

Water (Aqua), Isopropyl Alcohol, PEG-12 Dimethicone, Avena Sativa (Oat) Kernel Extract, Caprylyl glycol, Glycerin, Niacinamide, PPG-12/SMDI copolymer, Potassium Sorbate



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Purell Advanced Hand Sanitizer
ULTRA NOURISHING™ Foam

Drug Facts	
Active ingredient Ethyl alcohol 70% v/v	Purpose Antimicrobial
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Drug Facts (continued)
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*Based on a study comparing skin hydration over a 2-week period

DSP-0H-36



1200 mL
(40.5 FL OZ)
Reorder No. 5057



PURELL ADVANCED HAND SANITIZER ULTRA NOURISHING FOAM
alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:21749-856
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
PEG-12 DIMETHICONE (300 CST) (UNII: ZEL54N6W95)	
OAT (UNII: Z6J799EAJK)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
GLYCERIN (UNII: PDC6A3C0OX)	
NIACINAMIDE (UNII: 25X51I8RD4)	
PPG-12/SMDI COPOLYMER (UNII: 1BK9DDDD24E)	
POTASSIUM SORBATE (UNII: 1VPU26JZ Z4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21749-856-17	515 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/04/2023	
2	NDC:21749-856-40	1200 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/04/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	12/04/2023	

Labeler - GOJO Industries, Inc. (004162038)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	manufacture(21749-856) , pack(21749-856) , label(21749-856)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		088312414	manufacture(21749-856) , pack(21749-856) , label(21749-856)

Revised: 12/2024

GOJO Industries, Inc.