PURELL ADVANCED HAND SANITIZER ULTRA NOURISHING FOAMalcohol 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, Niaciniamide, PPG-12/SMDI copolymer, Potassium Sorbate



ADVANCED HAND **SANITIZER**

ULTRA NOURISHING™ Improves skin with repeated use*



Advanced Hand Sanitizer ULTRA NOURISHING™ Foam

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

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Drug Facts (continued)

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 Rub hands together briskly until dry

Inactive ingredients

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

*Based on a study comparing skin hydration over a 2-week period



1200 mL (40.5 FL 0Z)





Advanced Hand Sanitizer ULTRA NOURISHING™ Foam



ADVANCED

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DSP-OH-36

Drug Facts (continued)

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PURELL ADVANCED HAND SANITIZER ULTRA NOURISHING FOAM

alcohol liquid

Product Information

Product Type HUMAN OTC DRUG NDC:21749-856 Item Code (Source)

TOPICAL Route of Administration

Active Ingredient/Active Moiety

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

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: 1BK9DDD24E) POTASSIUM SORBATE (UNII: 1VPU26|ZZ4)

Packaging

	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
		NDC:21749- 856-17	515 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/04/2023	
		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: 3/2023 GOJO Industries, Inc.