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

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

PURELL Healthcare Advanced Hand Sanitizer ULTRA NOURISHING™ Foam

Active ingredient

Ethyl Alcohol 70% v/v

Purpose

Antimicrobial

Use

• 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, Canarium Luzonicum Gum Nonvolatiles, Caprylyl glycol, Citrus Aurantifolia (Lime) Peel Oil, Citrus Aurantium Bergamia (Bergamot) Oil, Citrus Aurantium Dulcis (Orange) Peel Oil, Citrus Limon (Lemon) Peel Oil, Glycerin, Isopropyl Myristate, Litsea Cubeba Fruit Oil, Mentha Arvensis Leaf Oil, Mentha Viridis (Spearmint) Leaf Oil, Niaciniamide, Pinus Silvestrus Leaf Oil, Pogostemon Cablin Oil, PPG-12/SMDI copolymer, Schinus Molle Oil, Tocopheryl Acetate, Potassium Sorbate

Purell®

ADVANCED

HAND SANITIZER

ULTRA NOURISHING™
Improves skin with repeated use

Purell®

Healthcare Advanced Hand Sanitizer ULTRA NOURISHING™ Foam

Drug Facts

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Drug Facts (continued)

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Patent Pending DSP-OH-36

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SAN
ES6

40.5 FL OZ
(1200 mL)
Reorder No. 6456

6456-640-B

PURELL HEALTHCARE ADVANCED HAND SANITIZER ULTRA NOURISHING FOAM

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:21749-854
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)	
PEG-12 DIMETHICONE (300 CST) (UNII: ZEL54N6W95)	
OAT (UNII: Z6J799EAJK)	
ELEMI (UNII: C13XI009KO)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
LIME OIL (UNII: UZH29XGA8G)	
BERGAMOT OIL (UNII: 39W1PKE3JI)	
ORANGE OIL (UNII: AKN3KSD11B)	
LEMON OIL (UNII: I9GRO824LL)	
GLYCERIN (UNII: PDC6A3C0OX)	
Isopropyl Myristate (UNII: 0RE8K4LNJS)	
LITSEA OIL (UNII: 2XIW34BN6O)	
MENTHA ARVENSIS LEAF OIL (UNII: 1AEY1M553N)	
SPEARMINT OIL (UNII: C3M81465G5)	
NIACINAMIDE (UNII: 25X51I8RD4)	
PINE NEEDLE OIL (PINUS SYLVESTRIS) (UNII: 5EXL5H740Y)	
PATCHOULI OIL (UNII: F3IN55X5PO)	
PPG-12/SMDI COPOLYMER (UNII: 1BK9DDD24E)	
SCHINUS MOLLE FRUIT OIL (UNII: 99O5U5NLK2)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:21749-854-53	535 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/30/2020	
2	NDC:21749-854-89	1200 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/15/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	10/15/2017	

Labeler - GOJO Industries, Inc. (004162038)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	manufacture(21749-854)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		088312414	manufacture(21749-854) , label(21749-854) , pack(21749-854)

Revised: 5/2020 GOJO Industries, Inc.