

PURELL ADVANCED HAND SANITIZER NATURALS 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 Advanced Hand Sanitizer Naturals Foam

Active ingredient

Ethyl alcohol 70% v/v

Purpose

Antimicrobial

Uses

- Hand sanitizer to help reduce bacteria on the skin

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
- Run hands together briskly until dry
- Children under 6 years of age should be supervised when using this product

Inactive ingredients

Water (Aqua), Isopropyl Alcohol, PEG-12 Dimethicone, Caprylyl Glycol, Citrus Aurantium Dulcis (Orange) Peel Oil, Glycerin, Lavandula Hybrida (Lavandin) Oil, Litsea Cubeba Fruit Oil, Pelargonium Graveolens (Geranium) Oil, Pogostemon Cablin Oil

NEW FOAM



Purell
ADVANCED
**HAND SANITIZER
FOAM**

NATURALS
MADE WITH PLANT BASED ALCOHOL

Kills 99.99% of Germs*
With Essential Oils
and Skin Conditioners

10 FL OZ (295 mL)

Purell
ADVANCED
**HAND SANITIZER
FOAM**

NATURALS

Foam feels good on hands, and 100% naturally renewable ethanol means you can feel good about using it, too.

- Formulated with 91% biobased ingredients
- Naturally fragrances
- Triclosan, paraben, and preservative free

Active ingredient
Ethyl Alcohol 70% v/v

*Kills 99.99% of most common germs that may cause illness.

Distributed by: GOJO Industries, Inc., Akron, OH 44316
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Made in USA with U.S. and foreign components.
DS9-041-08 3001-610-C01-P3

0 73852 129296

Drug Facts	
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alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:21749-875
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 (UNII: ZEL54N6W95)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	

ORANGE OIL, COLD PRESSED (UNII: AKN3KSD11B)	
GLYCERIN (UNII: PDC6A3C0OX)	
LAVANDIN OIL (UNII: 9RES347CKG)	
LITSEA OIL (UNII: 2XIW34BN6O)	
PELARGONIUM GRAVEOLENS FLOWER OIL (UNII: 3K0J1S7QGC)	
POGOSTEMON CABLIN LEAF OIL (UNII: F3IN55X5PO)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21749-875-10	295 mL in 1 PACKAGE; Type 0: Not a Combination Product	04/15/2021	
2	NDC:21749-875-50	500 mL in 1 PACKAGE; Type 0: Not a Combination Product	04/15/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	04/15/2021	

Labeler - GOJO Industries, Inc. (004162038)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	manufacture(21749-875) , pack(21749-875) , label(21749-875)

Revised: 4/2021

GOJO Industries, Inc.