PURELL GREEN CERTIFIED INSTANT HAND SANITIZER FOAM (ETHYL ALCOHOL)- 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 Green Certified Instant Hand Sanitizer 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

- Put enough product in your palm to cover hands and rub hands together briskly until dry
- Children under 6 years of age should be supervised when using PURELL

Inactive ingredients

Water (Aqua), Isopropyl Alcohol, PEG-10 Dimethicone, Glycerin, Hydroxyethyl Urea, Isopropyl Myristate, PEG-12 Dimethicone, Propylene Glycol, Tocopheryl Acetate

PURELL GREEN CERTIFIED INSTANT HAND SANITIZER FOAM (ETHYL ALCOHOL)

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:21749-987
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.70 mL in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	

PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21749-987-53	535 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	12/15/2010	
2	NDC:21749-987-51	550 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	12/15/2010	
3	NDC:21749-987-97	700 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/15/2010	
4	NDC:21749-987-10	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/15/2010	
5	NDC:21749-987-89	1200 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/15/2010	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	12/15/2010	

Labeler - GOJO Industries, Inc. (004162038)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		088312414	label(21749-987) , pack(21749-987) , manufacture(21749-987)

Revised: 10/2017 GOJO Industries, Inc.