PURELL ADVANCED INSTANT HAND SANITIZER- alcohol liquid
GOJO Industries, Inc.

PURELL Advanced Instant Hand Sanitizer Foam

Active ingredient

Ethyl Alcohol 70%

Purpose

Antimicrobial

Use

Hand sanitizer to help reduce bacteria on the skin that could cause disease

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.

Children under 6 years of age should be supervised when using this product.

Other information

Store below 110 ^{o}F (43 ^{o}C)

May discolor certain fabrics or surfaces

Inactive ingredients

Water (Aqua), Isopropyl Alcohol, PEG-12 Dimethicone, Caprylyl Glycol, Glycerin, Isopropyl Myristate, Tocopheryl Acetate, Fragrance (Parfum)

PURELL ADVANCED INSTANT HAND SANITIZER

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:21749-807
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)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
PEG-12 DIMETHICONE (300 CST) (UNII: ZEL54N6W95)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21749-807-31	45 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/11/2011	01/31/2024
2	NDC:21749-807-22	200 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/11/2011	12/13/2024
3	NDC:21749-807-23	222 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/11/2011	12/13/2024
4	NDC:21749-807-17	515 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/13/2022	
5	NDC:21749-807-53	535 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/11/2011	12/31/2024
6	NDC:21749-807-51	550 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/11/2011	
7	NDC:21749-807-33	1000 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/11/2011	
8	NDC:21749-807-89	1200 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/11/2011	
9	NDC:21749-807-97	700 mL in 1 PACKAGE; Type 0: Not a Combination Product	04/11/2011	
10	NDC:21749-807-30	300 mL in 1 PACKAGE; Type 0: Not a Combination Product	08/24/2020	12/13/2024
11	NDC:21749-807-50	189271 mL in 1 PACKAGE; Type 0: Not a Combination Product	08/24/2020	
12	NDC:21749-807-80	800 mL in 1 PACKAGE; Type 0: Not a Combination Product	02/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	04/11/2011	

Labeler - GOJO Industries, Inc. (004162038)

Registrant - GOJO Industries, Inc. (004162038)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	manufacture(21749-807)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		088312414	manufacture(21749-807) , label(21749-807) , pack(21749-807)

Revised: 1/2026 GOJO Industries, Inc.