PURELL ADVANCED HAND SANITIZER NATURALS GEL- alcohol gel
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 Gel

Active ingredient

Ethyl alcohol 70% v/v

Purpose

Antimicrobial

Use

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

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

Inactive Ingredients:

Water (Aqua), Isopropyl Alcohol, Caprylyl glycol, Citrus Aurantium Dulcis (Orange) Peel Oil, Glycerin, Isopropyl Myristate, Lavandula Hybrida (Lavandin) oil, Litsea Cubeba Fruit Oil, Pelargonium Graevolens (Geranium) oil, Pogostemon Cablin Oil, Tocopheryl Acetate, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aminomethyl Propanol


Purell
ADVANCED
HAND SANITIZER
NATURALS
MADE WITH PLANT BASED ALCOHOL
Kills 99.99% of Germs*
With Essential Oils
and Skin Conditioners
6.5 FL OZ (192 mL) 3265-640-CMR-F


Distributed by: GOJO Industries, Inc., Akron, OH 44309
Questions? Tel: 1-888-4-PURELL www.PURELL.com
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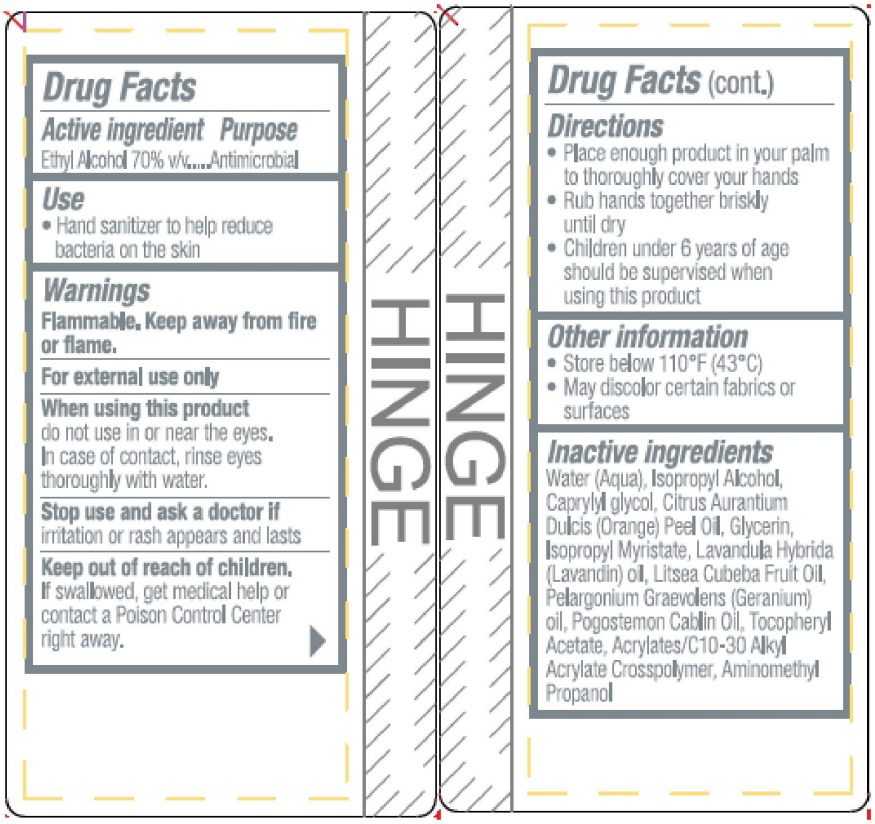
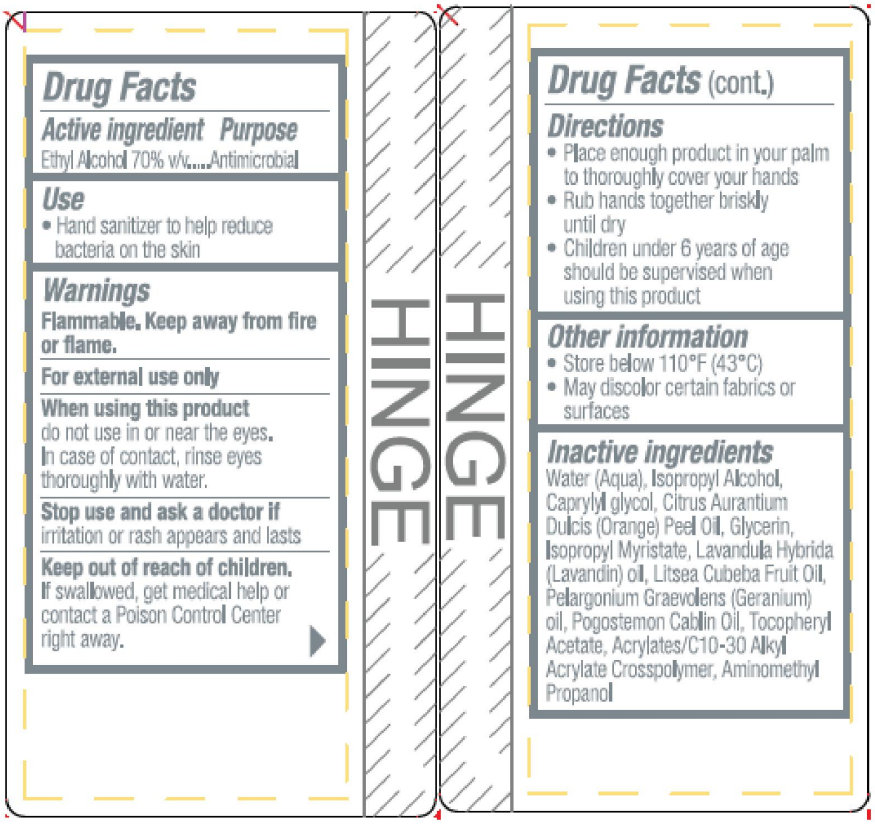
Made in the USA with U.S. and foreign components
3265-640-CMR-B
0 73852 29352 4
Purell
ADVANCED
HAND SANITIZER
NATURALS
100% naturally renewable ethanol supports well-being for your family and your world.
Ingredients are 93% naturally derived
Naturally fragranced
Triclosan, paraben, and preservative free
Active ingredient
Ethyl Alcohol 70% v/v
*Kills 99.99% of most common germs that may cause illness.


Drug Facts
Active ingredient Purpose
Ethyl Alcohol 70% v/v.....Antimicrobial
Use
• 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.
HINGE
HINGE
Drug Facts (cont.)
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°F (43°C)
• May discolor certain fabrics or surfaces
Inactive ingredients
Water (Aqua), Isopropyl Alcohol, Caprylyl glycol, Citrus Aurantium Dulcis (Orange) Peel Oil, Glycerin, Isopropyl Myristate, Lavandula Hybrida (Lavandin) oil, Litsea Cubeba Fruit Oil, Pelargonium Graevolens (Geranium) oil, Pogostemon Cablin Oil, Tocopheryl Acetate, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aminomethyl Propanol



PURELL ADVANCED HAND SANITIZER NATURALS GEL

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:21749-625
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)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
BITTER ORANGE OIL (UNII: 9TLV70SV6I)	
LAVANDIN OIL (UNII: 9RES347CKG)	
LITSEA OIL (UNII: 2XIW34BN6O)	
GERANIUM OIL, ALGERIAN TYPE (UNII: 5Q1I94P4WG)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
PATCHOULI OIL (UNII: F3IN55X5PO)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21749-625-01	30 mL in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2016	
2	NDC:21749-625-02	59 mL in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2016	
3	NDC:21749-625-04	118 mL in 1 PACKAGE; Type 0: Not a Combination Product	12/15/2019	
4	NDC:21749-625-08	236 mL in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2016	
5	NDC:21749-625-29	295 mL in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2016	
6	NDC:21749-625-12	355 mL in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2016	
7	NDC:21749-625-28	828 mL in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2016	
8	NDC:21749-625-10	1000 mL in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2016	
9	NDC:21749-625-24	710 mL in 1 PACKAGE; Type 0: Not a Combination Product	08/01/2021	
10	NDC:21749-	192 mL in 1 PACKAGE; Type 0: Not a Combination	07/01/2021	

10	625-65	Product	07/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	01/01/2016	

Labeler - GOJO Industries, Inc. (004162038)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	manufacture(21749-625)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		088312414	manufacture(21749-625) , label(21749-625) , pack(21749-625)

Revised: 5/2021 GOJO Industries, Inc.