

PURELL ADVANCED INSTANT HAND SANITIZER- 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

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

Inactive ingredients

Water (Aqua), Isopropyl Alcohol, Caprylyl Glycol, Glycerin, Isopropyl Myristate, Tocopheryl Acetate, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aminomethyl Propanol, Fragrance (Parfum)



ADVANCED
INSTANT **HAND
SANITIZER**
with MOISTURIZERS
and VITAMIN E

AVANZADOS
ANTISÉPTICO **INSTANTÁNEO
PARA LAS MANOS**
con HUMECTANTES
y VITAMINA E

Distributed by / Distribuido por:
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Made in U.S.A., Hecho en los E.E.U.U.
Patent Pending / Patente Pendiente
SDA-36-1301 SDA-34-962



NDC 21749-700-10

Drug Facts

Active ingredient
Ethyl alcohol 70% w/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 • 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 • Do not store above 110°F (43°C) • May discolor certain fabrics or surfaces

Inactive ingredients

Water (Aqua), Isopropyl Alcohol, Caprylyl Glycol, Glycerin, Isopropyl Myristate, Tocopheryl Acetate, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aminomethyl Propanol, Fragrance (Parfum)

* Meets Food Code Hand Sanitizers criteria (Section 2-301.16) published by the FDA / * Cumple Con Los criterios Del Código De Alimentos Para Esterilizadores De Manos (Sección 2-301.16)

Datos Farmacológicos (cont'd)

Ingrediente activo
Alcohol etílico 70% w/v.....

Propósito
Antimicrobiano

Usos • Antiséptico para las manos, empleado para disminuir la cantidad de bacterias en la piel que pueden causar enfermedades • Recomendado para uso reiterado

Advertencias

Inflamable. Mantener alejado del fuego o las llamas.

Sólo para uso externo

Al utilizar este producto, evitar el contacto con los ojos o con la zona alrededor de los ojos. En caso de contacto, enjuagar completamente los ojos con agua.

Dejar de usar el producto y consultar a un médico si aparece y persiste una irritación o erupción cutánea

Mantener fuera del alcance de los niños. En caso de ingestión, de inmediato acudir a un médico o ponerse en contacto con un centro para el control de tóxicos.

Modo de uso • Poner suficiente cantidad del producto en la palma de la mano para cubrir las manos completamente • Frotarse las manos vigorosamente hasta que se sequen • El empleo de este producto por niños menores de seis años deberá ser supervisado

Información adicional • Almacene a temperaturas inferiores a 43°C (110°F) • Puede decolorar ciertas telas o superficies

Ingredientes inactivos

Agua, Alcohol isopropílico, Caprilil glicol, Glicerina, Miristato de isopropilo, Acetato de tocoferilo, Copolímero de acrilatos/C10-30 alquil acrilato, Aminometil propanol, Fragancia

U.S. Pat. 6,619,512
U.S. Pat. 6,877,642 • U.S. Pat. D431,404
U.S. Pat. 6,216,916 • U.S. Pat. D432,547

1 L (33.8 US/ÉU FL OZ)

2156

PURELL ADVANCED INSTANT HAND SANITIZER

alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:21749-700
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)	

.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)

Caprylyl Glycol (UNII: 00YIU5438U)

AMINOMETHYLPROPANOL (UNII: LU49E6626Q)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21749-700-02	24 in 1 CASE	05/10/2011	
1		59 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:21749-700-10	8 in 1 CASE	05/10/2011	
2		1000 mL in 1 BAG; Type 0: Not a Combination Product		
3	NDC:21749-700-89	4 in 1 CASE	05/10/2011	
3		1200 mL in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:21749-700-20	4 in 1 CASE	05/10/2011	
4		2000 mL in 1 BAG; Type 0: Not a Combination Product		
5	NDC:21749-700-67	4 in 1 CASE	05/10/2011	
5		2000 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product		
6	NDC:21749-700-04	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/10/2011	
7	NDC:21749-700-80	800 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/10/2011	
8	NDC:21749-700-12	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/10/2011	
9	NDC:21749-700-07	222 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/10/2011	
10	NDC:21749-700-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/10/2011	
11	NDC:21749-700-59	590 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/10/2011	
12	NDC:21749-700-15	15 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/10/2011	
13	NDC:21749-700-60	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/10/2011	
14	NDC:21749-700-47	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/10/2011	
15	NDC:21749-700-03	89 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/10/2011	
16	NDC:21749-700-08	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/10/2011	
17	NDC:21749-700-00	1.2 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2016	
18	NDC:21749-700-45	450 mL in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2016	
19	NDC:21749-700-29	295 mL in 1 PACKAGE; Type 0: Not a Combination Product	09/28/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	05/10/2011	

Labeler - GOJO Industries, Inc. (004162038)

Registrant - GOJO Industries, Inc. (004162038)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	manufacture(21749-700)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		088312414	manufacture(21749-700) , label(21749-700) , pack(21749-700)

Establishment

Name	Address	ID/FEI	Business Operations
Travis Association for the Blind		026032268	pack(21749-700)

Revised: 12/2019

GOJO Industries, Inc.