# PURELL HEALTHCARE ADVANCED HAND SANITIZER 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.

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#### **PURELL Healthcare Advanced Hand Sanitizer Gel**

#### 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

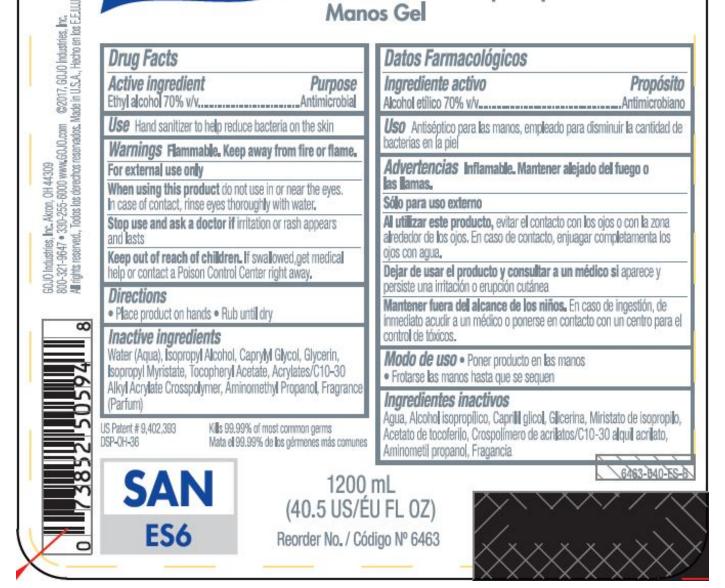
- Place product on hands
- Rub until dry

#### Inactive ingredients

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

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### Healthcare Advanced Hand Sanitizer Gel Avanzados Antiséptico para Manos Gel



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Drug Facts	Datos Farmacológicos
Active ingredient Purpose Ethyl alcohol 70% v/v	Ingrediente activo Propósito Alcohol etílico 70% v/vAntimicrobiano
Use Hand sanitizer to help reduce bacteria on the skin	<b>Uso</b> Antiséptico para las manos, empleado para disminuir la cantidad de bacterias en la piel
Warnings Flammable, Keep away from fire or flame,	Advertencias Inflamable. Mantener alejado del fuego o
For external use only When using this product do not use in or near the eyes.	las llamas.
In case of contact, rinse eyes thoroughly with water.	Sólo para uso externo
Stop use and ask a doctor if irritation or rash appears and lasts	Al utilizar este producto, evitar el contacto con los ojos o con la zona alrededor de los ojos. En caso de contacto, enjuagar completamenta los
Keep out of reach of children. If swallowed, get medical	ojos con agua.
help or contact a Poison Control Center right away. Directions	Dejar de usar el producto y consulitar a un médico si aparece y persiste una initación o erupción cutánea.
Place product on hands  Rub until dry	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
Inactive ingredients	control de tóxicos.
Water (Aqua), Isopropyl Alcohol, Caprylyl Glycol, Glycerin, Isopropyl Myristate, Tocopheryl Acetate, Acrylates/C10-30	Modo de uso • Poner producto en las manos • Frotarse las manos hasta que se seguen
Alkyl Acrylate Crosspolymer, Aminomethyl Propanol, Fragrance (Parfum)	Ingredientes inactivos
(Partoin) US Patent # 9,402,393 Kills 99,99% of most common germs	Agua, Alcohol isopropílico, Caprilil glicol, Glicerina, Miristato de isopropilo, Acetato de tocoferilo, Crospolímero de acrilatos/C10-30 alquil acritato,
DSP-0H-36 Mata el 99.99% de los gérmenes más comunes	Acetato de tocoferilo, Crospolimero de acritatos/C10-30 alquit acritato, Aminometil propanol, Fragancia
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### PURELL HEALTHCARE ADVANCED HAND SANITIZER GEL

alcohol gel

Product Information				
Product Type	HUMAN OTC DRUG	ltem Co	de (Source)	NDC:21749-716
Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
Ingredient Name			<b>Basis of Strengt</b>	h 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)		
GLYCERIN (UNII: PDC6A3C0OX)		
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)		
.ALPHATOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)		
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)		

### Packaging

#	ltem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21749- 716-01	29 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/15/2017	
2	NDC:21749- 716-02	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/15/2017	
3	NDC:21749- 716-04	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/15/2017	
4	NDC:21749- 716-08	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/15/2017	
5	NDC:21749- 716-12	354 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/15/2017	
6	NDC:21749- 716-89	1200 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/15/2017	
7	NDC:21749- 716-20	2000 mL in 1 PACKAGE; Type 0: Not a Combination Product	10/15/2017	

## **Marketing Information**

Marketing	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date
OTC monograph not final	part333E	10/15/2017	

Labeler - GOJO Industries, Inc. (004162038)

Registrant - GOJO Industries, Inc. (004162038)

Establishment				
Name	Address	ID/FEI	<b>Business Operations</b>	
GOJO Industries, Inc.		036424534	MANUFACTURE(21749-716)	
Establishment				

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		088312414	MANUFACTURE(21749-716) , label(21749-716) , pack(21749-716)

Revised: 12/2022