

**PURELL HEALTHCARE ADVANCED HAND SANITIZER GEL- alcohol gel**  
**GOJO Industries, Inc.**

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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)



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

GOJO Industries, Inc. Akron, OH 44309  
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**Drug Facts**

**Active ingredient** **Purpose**  
Ethyl alcohol 70% v/v.....Antimicrobial

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and lasts

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help or contact a Poison Control Center right away.

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Alkyl Acrylate Crosspolymer, Aminomethyl Propanol, Fragrance  
(Parfum)

US Patent # 9,402,393  
DSP-OH-36

Kills 99.99% of most common germs  
Mata el 99.99% de los gérmenes más comunes

**SAN**  
**ES6**

1200 mL  
(40.5 US/ÉU FL OZ)  
Reorder No. / Código Nº 6463

6463-040-ES-6

**Datos Farmacológicos**

**Ingrediente activo** **Propósito**  
Alcohol etílico 70% v/v.....Antimicrobiano

**Uso** Antiséptico para las manos, empleado para disminuir la cantidad de  
bacterias en la piel

**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 producto en las manos

- Frotarse las manos hasta que se sequen

**Ingredientes inactivos**

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

**PURELL HEALTHCARE ADVANCED HAND SANITIZER GEL**

alcohol gel

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:21749-716
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>ALCOHOL</b> (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.7 mL in 1 mL

## Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>ISOPROPYL ALCOHOL</b> (UNII: ND2M416302)	
<b>CAPRYLYL GLYCOL</b> (UNII: 00YIU5438U)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>ISOPROPYL MYRISTATE</b> (UNII: ORE8K4LNJS)	
<b>.ALPHA.-TOCOPHEROL ACETATE, D-</b> (UNII: A7E6112E4N)	
<b>AMINOMETHYLPROPANOL</b> (UNII: LU49E6626Q)	

## Packaging

#	Item 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	03/31/2022
2	NDC:21749-716-02	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/15/2017	05/31/2022
3	NDC:21749-716-04	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/15/2017	12/11/2024
4	NDC:21749-716-08	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/15/2017	12/11/2024
5	NDC:21749-716-12	354 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/15/2017	12/11/2024
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	12/11/2024

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	10/15/2017	

**Labeler** - GOJO Industries, Inc. (004162038)

**Registrant** - GOJO Industries, Inc. (004162038)

## Establishment

Name	Address	ID/FEI	Business Operations
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)

