

**PURELL PROFESSIONAL ADVANCED HAND SANITIZER FRAGRANCE FREE 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 Professional Advanced Hand Sanitizer Fragrance Free Gel

Active ingredient

Ethyl alcohol 70% v/v

Purpose

Antimicrobial

Uses

- Hand sanitizer to help reduce bacteria on the skin that 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 product on hands
- Run until dry

Inactive ingredients

Water (Aqua), Isopropyl Alcohol, PEG-12 Dimethicone, Caprylyl Glycol, Glycerin, Isopropyl Myristate, Tocopheryl Acetate, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aminomethyl Propanol



Professional Advanced Hand Sanitizer Fragrance Free Gel

Avanzados Antiséptico para Manos
Gel sin Fragancia

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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.

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

US Patent # 9,402,393

DSP-OH-36

SAN
ES4

1200 mL
(40.5 US/ÉU FL OZ)
Reorder No. / Código Nº 5060



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, Crospolímico de acrilatos/C10-30 alquil acrilato,
Aminometil propanol

PURELL PROFESSIONAL ADVANCED HAND SANITIZER FRAGRANCE FREE GEL

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:21749-717
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.7 mL in 1 mL
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Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21749-717-08	236 mL in 1 PACKAGE; Type 0: Not a Combination Product	10/15/2017	
2	NDC:21749-717-12	354 mL in 1 PACKAGE; Type 0: Not a Combination Product	10/15/2017	
3	NDC:21749-717-10	1000 mL in 1 PACKAGE; Type 0: Not a Combination Product	10/15/2017	
4	NDC:21749-717-89	1200 mL in 1 PACKAGE; Type 0: Not a Combination Product	10/15/2017	
5	NDC:21749-717-20	2000 mL in 1 PACKAGE; Type 0: Not a Combination Product	10/15/2017	

Marketing Information

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

Labeler - GOJO Industries, Inc. (004162038)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	MANUFACTURE(21749-717)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		088312414	manufacture(21749-717) , label(21749-717) , pack(21749-717)