

**PURELL PROFESSIONAL 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 Professional 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)



# Professional Advanced Hand Sanitizer Gel

GOJO Industries, Inc. Akron, OH 44309  
800-321-9647 • 330-255-8000 www.GOJO.com  
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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. ▶

US Patent # 9,402,393

Kills 99.99% of most common germs

DSP-OH-36

### Drug Facts (continued)

#### 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)



1000 mL  
(33.8 US/ÉU FL OZ)

Reorder No. / Código N° 2162

2162-640-F



# Professional Advanced Hand Sanitizer Gel

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|  |                                 |
|--|---------------------------------|
| <b>Drug Facts</b>  |                                 |
| <b>Active ingredient</b><br>Ethyl alcohol 70% v/v.....   | <b>Purpose</b><br>Antimicrobial |
| <b>Use</b> Hand sanitizer to help reduce bacteria on the skin  |                                 |
| <b>Warnings</b> Flammable. Keep away from fire or flame.   |                                 |
| <b>For external use only</b>   |                                 |
| <b>When using this product</b> do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water. |                                 |
| <b>Stop use and ask a doctor</b> if irritation or rash appears and lasts   |                                 |
| <b>Keep out of reach of children.</b> If swallowed, get medical help or contact a Poison Control Center right away.  |                                 |

US Patent # 9,402,393 DSP-OH-36  
Kills 99.99% of most common germs

|  |  |
|--|--|
| <b>Drug Facts</b> (continued)  |  |
| <b>Directions</b><br>• Place product on hands • Rub until dry  |  |
| <b>Inactive ingredients</b><br>Water (Aqua), Isopropyl Alcohol, Caprylyl Glycol, Glycerin, Isopropyl Myristate, Tocopheryl Acetate, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aminomethyl Propanol, Fragrance (Parfum) |  |



1000 mL  
(33.8 US/ÉU FL OZ)  
Reorder No. / Código N° 2162

2162-640-F

## PURELL PROFESSIONAL ADVANCED HAND SANITIZER GEL alcohol gel

|                                |                |                           |               |
|--------------------------------|----------------|---------------------------|---------------|
| <b>Product Information</b>     |                |                           |               |
| <b>Product Type</b>            | HUMAN OTC DRUG | <b>Item Code (Source)</b> | NDC:21749-715 |
| <b>Route of Administration</b> | TOPICAL        |                           |               |

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

|                                      |                 |  |  |
|--------------------------------------|-----------------|--|--|
| <b>Inactive Ingredients</b>          |                 |  |  |
| <b>Ingredient Name</b>               | <b>Strength</b> |  |  |
| WATER (UNII: 059QF0KO0R)             |                 |  |  |
| ISOPROPYL ALCOHOL (UNII: ND2M416302) |                 |  |  |

|  |  |
|--|--|
| <b>CAPRYLYL GLYCOL</b> (UNII: 00YIU5438U)                |  |
| <b>GLYCERIN</b> (UNII: PDC6A3C0OX)                       |  |
| <b>ISOPROPYL MYRISTATE</b> (UNII: 0RE8K4LNJS)            |  |
| <b>.ALPHA.-TOCOPHEROL ACETATE, D-</b> (UNII: A7E6112E4N) |  |
| <b>AMINOMETHYLPROPANOL</b> (UNII: LU49E6626Q)            |  |

| <b>Packaging</b> |                  |   |                      |                    |
|------------------|------------------|---|----------------------|--------------------|
| #                | Item Code        | Package Description                                     | Marketing Start Date | Marketing End Date |
| 1                | NDC:21749-715-08 | 236 mL in 1 BOTTLE; Type 0: Not a Combination Product   | 10/15/2017           |                    |
| 2                | NDC:21749-715-12 | 354 mL in 1 BOTTLE; Type 0: Not a Combination Product   | 10/15/2017           |                    |
| 3                | NDC:21749-715-10 | 1000 mL in 1 BOTTLE; Type 0: Not a Combination Product  | 10/15/2017           |                    |
| 4                | NDC:21749-715-89 | 1200 mL in 1 BOTTLE; Type 0: Not a Combination Product  | 10/15/2017           |                    |
| 5                | NDC:21749-715-20 | 2000 mL in 1 PACKAGE; Type 0: Not a Combination Product | 10/15/2017           |                    |

| <b>Marketing Information</b> |  |                      |                    |
|------------------------------|--|----------------------|--------------------|
| 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)

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

| <b>Establishment</b>  |         |           |                        |
|-----------------------|---------|-----------|------------------------|
| Name                  | Address | ID/FEI    | Business Operations    |
| GOJO Industries, Inc. |         | 036424534 | MANUFACTURE(21749-715) |

| <b>Establishment</b>  |         |           |   |
|-----------------------|---------|-----------|---|
| Name                  | Address | ID/FEI    | Business Operations   |
| GOJO Industries, Inc. |         | 088312414 | MANUFACTURE(21749-715) , label(21749-715) , pack(21749-715) |