

PURELL HAND SANITIZING WIPES- benzalkonium chloride cloth
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 Hand Sanitizing Wipes

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antimicrobial

Uses

- Hand sanitizer to help reduce bacteria on the skin

Warnings

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

- Wet hands thoroughly with product and allow to dry
- Children under 6 years of age should be supervised when using PURELL

Inactive ingredients

Water (Aqua), Decyl Glucoside, Glycerin, Fragrance(Parfum), Phenoxyethanol



HAND SANITIZING WIPES

KILLS 99.99% OF MOST COMMON GERMS

**Dermatologist Tested
Paraben Free**

Distributed by:
GOJO Industries, Inc.
Akron, OH 44309
800-321-9647 • 330-255-6000
www.GOJO.com
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Nonfood Compounds
Program Listed E3 145050



Open Wipes Refill Pouch



Locate tear-notch on side of pouch.



Tear straight across to open.
Do not remove wipes roll from pouch.



Pull first wipe from center of roll up through opening.

Place in Dispensing Unit



Thread first wipe through dispensing nozzle in lid of floor stand.

- OR -



Thread first wipe through dispensing nozzle in top of wall mounted dispenser.

Reorder No. 9115

1500 Wet Wipes • 5 in. x 8 in. (12.7 cm x 20.3 cm)

9115-952



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benzalkonium chloride cloth

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:21749-368 |
| Route of Administration | TOPICAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|-----------------------|-------------------|
| BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y) | BENZALKONIUM CHLORIDE | 0.13 mg in 100 mL |

Inactive Ingredients

| Ingredient Name | Strength |
|---|----------|
| WATER (UNII: 059QF0KO0R) | |
| DECYL GLUCOSIDE (UNII: Z17H97EA6Y) | |
| GLYCERIN (UNII: PDC6A3C0OX) | |
| PHENOXYETHANOL (UNII: HIE492ZZ3T) | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:21749-368-35 | 35 in 1 PACKAGE | 03/14/2012 | 07/01/2022 |
| 1 | | 150 mL in 1 PACKAGE; Type 0: Not a Combination Product | | |
| 2 | NDC:21749-368-10 | 100 in 1 PACKAGE | 03/14/2012 | |
| 2 | | 245 mL in 1 PACKAGE; Type 0: Not a Combination Product | | |
| 3 | NDC:21749-368-27 | 270 in 1 PACKAGE | 03/14/2012 | |
| 3 | | 667 mL in 1 PACKAGE; Type 0: Not a Combination Product | | |
| 4 | NDC:21749-368-12 | 1200 in 1 PACKAGE | 03/14/2012 | |
| 4 | | 2567 mL in 1 PACKAGE; Type 0: Not a Combination Product | | |
| 5 | NDC:21749-368-15 | 1500 in 1 PACKAGE | 03/14/2012 | |
| 5 | | 2674 mL in 1 PACKAGE; Type 0: Not a Combination Product | | |
| 6 | NDC:21749-368-17 | 1700 in 1 PACKAGE | 05/01/2018 | |
| 6 | | 2273 mL in 1 PACKAGE; Type 0: Not a Combination Product | | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|-------------------------|--|----------------------|--------------------|
| OTC monograph not final | part333E | 03/14/2012 | |

Labeler - GOJO Industries, Inc. (004162038)

Establishment

| Name | Address | ID/FEI | Business Operations |
|-----------------------|---------|-----------|------------------------|
| GOJO Industries, Inc. | | 036424534 | MANUFACTURE(21749-368) |

