

**PURELL HAND SANITIZING WIPES- benzalkonium chloride cloth**  
**GOJO Industries, Inc.**

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**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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## Drug Facts

### Active ingredient

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**Questions?** Call 1-800-321-9647 • www.GOJO.com



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

## PURELL HAND SANITIZING WIPES

benzalkonium chloride cloth

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:21749-368
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

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

## Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>DECYL GLUCOSIDE</b> (UNII: Z17H97EA6Y)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>PHENOXYETHANOL</b> (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	09/30/2015
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 Drug	505G(a)(3)	03/14/2012	

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

## Establishment

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

