

**PURELL HAND SANITIZING WIPES FRESH CITRUS SCENT- benzalkonium chloride  
wipe liquid  
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 Hand Sanitizing Wipes Fresh Citrus Scent**

**Active ingredient**

Benzalkonium Chloride 0.13%

**Purpose**

Antimicrobial

**Use**

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

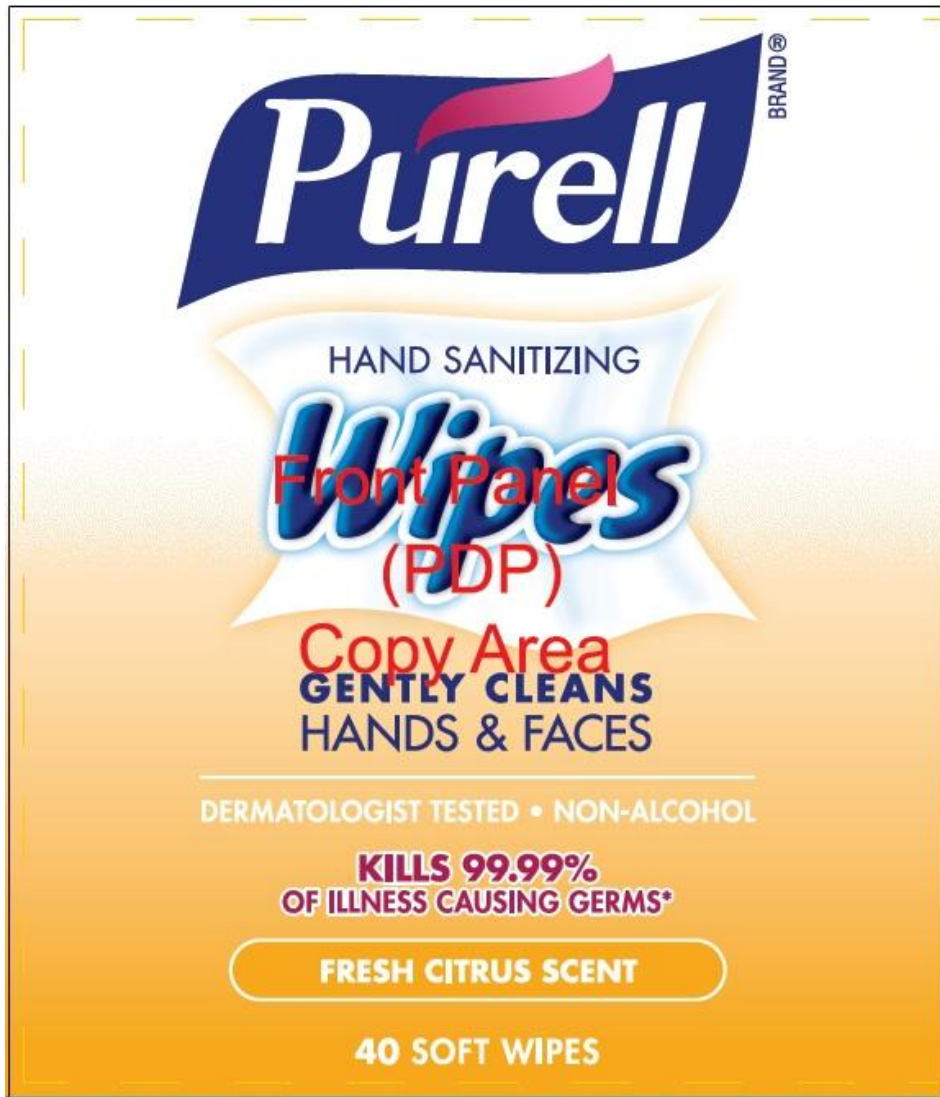
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

**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), Methylparaben, Phenoxyethanol, Propylparaben



<b>Drug Facts</b>	
<b>Active ingredient</b>	<b>Purpose</b>
Benzalkonium Chloride 0.13%	Antimicrobial
<b>Use</b>	
Hand sanitizer to help reduce bacteria on the skin	
<b>Warnings</b>	
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 the reach of children. If swallowed, get medical help or contact Poison Control Center right away.	
<b>Directions</b>	
<ul style="list-style-type: none"> <li>Wet hands thoroughly with product and allow to dry</li> <li>Discard wipe in trash receptacle after use. Do not flush.</li> <li>Children under 6 years of age should be supervised when using PURELL.</li> </ul>	
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**PURELL HAND SANITIZING WIPES FRESH CITRUS SCENT**  
benzalkonium chloride wipe liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:21749-358
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.00013

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
GLYCERIN (UNII: PDC6A3C0OX)	

METHYL PARABEN (UNII: A218C7H9T)

PHENOXYETHANOL (UNII: HIE492ZZ3T)

PROPYL PARABEN (UNII: Z8IX2SC1OH)

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21749-358-40	40 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	08/01/2014	

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

Revised: 8/2014

GOJO Industries, Inc.