

**PURELL SF607 INSTANT HAND SANITIZING- benzalkonium chloride 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 SF607 Instant Hand Sanitizing Foam**

**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**

- Place enough product in your palm to thoroughly cover your hands
- Rub hands together briskly until dry
- Children under 6 years of age should be supervised when using this product.

Water (Aqua), Decyl Glucoside, Glycerin, Methylparaben, Phenoxyethanol, Propylparaben

NDC 21749-607-89



SF607™  
INSTANT HAND  
SANITIZING FOAM

gojo 8802

Distributed by: GOJO Industries, Inc.  
Akron, OH 44309  
Questions? Tel: 800-321-9647 •  
330-255-6000 www.GOJO.com  
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Not for Retail Sale

1200 mL (40.5 US/ÉU FL OZ)



**Drug Facts**

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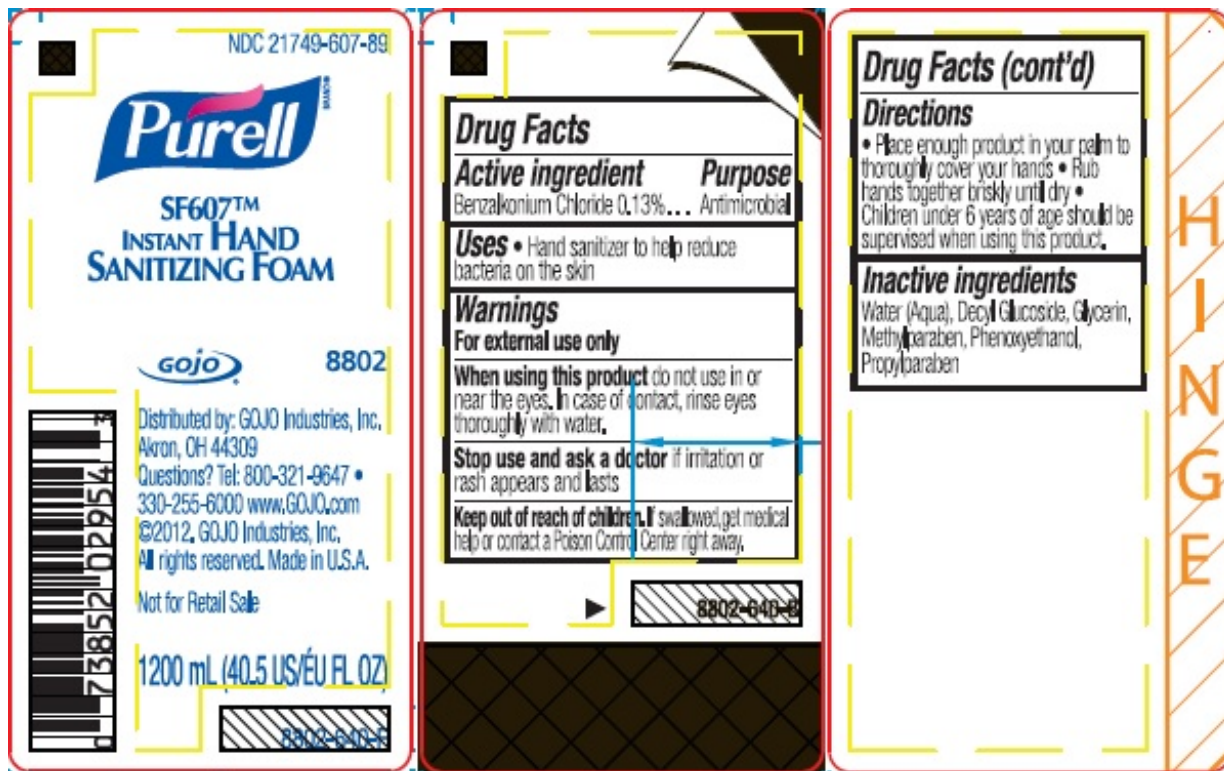
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## PURELL SF607 INSTANT HAND SANITIZING

benzalkonium chloride liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:21749-607
<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>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)	
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21749-607-31	45 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/03/2011	01/01/2024

2	NDC:21749-607-53	535 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/03/2011	01/01/2024
3	NDC:21749-607-89	1200 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/03/2011	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	01/03/2011	

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

## Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	manufacture(21749-607)

## Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		088312414	label(21749-607) , pack(21749-607)

## Establishment

Name	Address	ID/FEI	Business Operations
Travis Association for the Blind		026032268	label(21749-607)

Revised: 2/2023

GOJO Industries, Inc.