

**HAND SANITIZER- benzalkonium chloride 0.1%w/w liquid  
Georgia Pacific Consumer Products**

*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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**Alcohol Free Hand Sanitizer  
148.000/148AA**

**Claims**

enMotion

Foam Hand Sanitizer

with Moisturizers

Alcohol Free

Fragrance Free SKU 42338

**Active ingredients**

Benzalkonium Chloride 0.1% w/w

**Purpose**

Antiseptic

**Uses**

- To decrease bacteria on skin that could cause disease
- Recommended for repeated use

**Warnings**

**for external use only**

**When using this product**

do not use in or near eyes

**Discontinue use**

If irritation or redness develops. If condition persists for more than 72 hours, consult a physician

**Keep out of reach of children**

If swallowed, seek medical attention or call a poison control center immediately.

**Directions**

- Wet hands thoroughly with product and allow to dry without wiping.

**Inactive ingredients**

water, decyl glucoside, PEG-12 Dimethicone, Aloe Barbadensis Leaf Juice, Propylene Glycol, Glycerin, Panthenol Dipotassium Phosphate, Potassium Phosphate

**Adverse Reaction**

Manufacture for

Georgia-Pacific Consumer Products LP. Atlanta, GA 30303

Questions? Call 1-866-HELLOGP (435-5647)

or visit us online at [www.gppro.com](http://www.gppro.com)

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**Principal display panel**

Foam Hand Sanitizer

with Moisturizers

Alcohol Free

1000 mL (33.8 FL OZ)

42338-V2PRDRevA  
L0017235FA

**Foam Hand Sanitizer**  
**with Moisturizers**  
**Alcohol Free**

1000 mL (33.8 FL OZ)

## HAND SANITIZER

benzalkonium chloride 0.1%w/w liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:54622-148
<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	1 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>Water</b> (UNII: 059QF0KO0R)	
<b>DECYL GLUCOSIDE</b> (UNII: Z17H97EA6Y)	
<b>PEG-12 DIMETHICONE</b> (UNII: ZEL54N6W95)	

<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>PANTHENOL</b> (UNII: WW9CM0067Z)	
<b>DIBASIC POTASSIUM PHOSPHATE</b> (UNII: CI71S98N1Z)	
<b>POTASSIUM PHOSPHATE, UNSPECIFIED FORM</b> (UNII: B7862WZ 632)	

<b>Packaging</b>				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54622-148-20	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/01/2017	

<b>Marketing Information</b>			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	12/01/2017	

**Labeler** - Georgia Pacific Consumer Products (806142217)

**Registrant** - Vi-Jon, LLC (790752542)

<b>Establishment</b>			
Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(54622-148)

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Vi-Jon, LLC		790752542	manufacture(54622-148)