

# NOVAGEL ALCOHOL-FREE FOAMING HAND SANITIZER- benzalkonium chloride hand sanitizer liquid

Zhejiang Hongshiliang Health Technology Co Ltd

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.

## Principle Display Panel

**NOVA** Gel

**Drug Facts**

<b>Active Ingredient</b>	<b>Purpose</b>
Benzalkonium chloride 0.1% -----	Antimicrobial

**Use** • To decrease 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** • Pump a small amount of foam onto palm of hand • Rub thoroughly over all surfaces of both hands • Rub hands together briskly until dry

**Inactive ingredients** Water, Glycerin, Sodium Lauryl Sulfate, Fragrance

NDC: 78360-001-01

Distributed by Plasmadent, Inc. 3609 Mojave Ct, Suite E, Columbia, MO 65202

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**NOVA** Gel

**Alcohol-free**  
Foaming Hand Sanitizer

Lavender Scent

Leaves Hands Feeling Soft  
2 FL OZ (59.1 mL)

## Active Ingredient

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## Use

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Keep out of reach of children

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### Directions

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### Inactive ingredients

Inactive ingredients- water, glycerin, sodium lauryl sulfate, fragrance

### Dosage and Administration Section

Benzalkonium Chloride 0.1%

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### Purpose Section

Purpose

Antimicrobial

## NOVAGEL ALCOHOL-FREE FOAMING HAND SANITIZER

benzalkonium chloride hand sanitizer liquid

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78360-001
Route of Administration	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.1 g in 100 g

### Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	97.1 g in 100 g
GLYCERIN (UNII: PDC6A3C0OX)	2 g in 100 g
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	0.5 g in 100 g

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78360-001-01	59.15 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/26/2020	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/26/2020	

**Labeler** - Zhejiang Hongshiliang Health Technology Co Ltd (553365550)

Revised: 4/2020

Zhejiang Hongshiliang Health Technology Co Ltd