

**WOODWARDS HANDCLENS FOAMING SANITIZER COLOR AND FRAGRANCE FREE-
benzalkonium chloride liquid
Pacific World Corporation**

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.

Woodward's HandClens® Foaming Sanitizer & Lotion Color & Fragrance Free

Drug Facts

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antiseptic Sanitizer

Uses

- When soap and water are not available.
- Kills germs and helps prevent infections in minor cuts and abrasions.

Warnings

- ***Do not use*** in eyes, ears or mouth. In case of eye contact, flush with water immediately.
- ***Stop use and ask a doctor*** if rash or irritation occurs or if condition persists.
- ***Keep out of reach of children under 2 years of age.***

Directions

- Apply as needed and rub into skin.
- No rinsing necessary.

Inactive Ingredients

Water, Didecyldimonium Chloride, Methylparaben, Propylparaben, Propylene Glycol, Diazolidinyl Urea, Allantoin, Cocamidopropylamine Oxide, Cetrimonium Chloride, Cocamidopropyl Betaine, Triethanolamine, Citric Acid

PRINCIPAL DISPLAY PANEL - 236 mL Bottle Label

Color-Free
Fragrance-Free

**2 *in* 1
Sanitizer & Lotion**

**WOODWARD'S
HandClens®**

Foaming Sanitizer & Lotion

Alcohol-free • Non-Flammable

Softens Hands with Each Use!

***Kills 99.99%
of Germs***

8 FL OZ (236 mL)

Color-Free
Fragrance-Free

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Drug Facts **3X** more applications

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Manufactured for © Pacific World Corporation, Aliso Viejo, CA 92656 40-12091-0B1
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Made in USA
 800-780-6999
 www.HandClens.com
 U.S. Patent Nos. 5,661,170; 5,827,870.

WOODWARDS HANDCLENES FOAMING SANITIZER COLOR AND FRAGRANCE FREE

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:60193-201
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
DIDECYLDIMONIUM CHLORIDE (UNII: JXN40O9Y9B)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
ALLANTOIN (UNII: 344S277G0Z)	
COCAMIDOPROPYLAMINE OXIDE (UNII: M4SL82J7HK)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
TROLAMINE (UNII: 9O3K93S3TK)	
ALCOHOL (UNII: 3K9958V90M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date

1	NDC:60193-201-01	236 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product		
2	NDC:60193-201-03	1000 mL in 1 CARTRIDGE; Type 0: Not a Combination Product		
3	NDC:60193-201-05	1800 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product		
4	NDC:60193-201-07	50 mL in 1 BOTTLE, PUMP; 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	part333A	06/03/2013	

Labeler - Pacific World Corporation (089693097)

Revised: 10/2014

Pacific World Corporation