WOODWARDS HANDCLENS FOAMING SANITIZER- 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

Drug Facts

Active ingredient

Benzalkonium Chloride, 0.13%

Purpose

Antiseptic Hand Sanitizer

Use

for hand sanitizing to decrease bacteria on the skin

Warnings

- ***For external use only***

When using this product

- Avoid contact with eyes. In case of eye contact, flush with water
- Stop use and ask a doctor if rash or irritation develops and lasts

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

Directions

- Use enough foam to cover your hands
- Rub hands together briskly until dry

Other information

- Do not store above 110°F (40°C)
- You may report questions or adverse reaction from using this product to 1(541)-476-3178, 8 am to 5 pm (M–F), PST

Inactive ingredients

Water, Cocamidopropylamine Oxide, Didecyldimonium Chloride, Allantoin, Propylene Glycol, Cetrimonium Chloride, Cocamidopropyl Betaine, Sodium Hydroxide, Diazolidinyl Urea, Methylparaben, Propylparaben, Fragrance, Ext. Violet 2 (CI 60730), Green 5 (CI 61570).

PRINCIPAL DISPLAY PANEL - 50 mL Bottle Label

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

1.7 FL OZ (50 mL)

WOODWARDS HANDCLENS FOAMING SANITIZER

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:60193-202
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)	
D&C GREEN NO. 5 (UNII: 8J6RDU8L9X)	
EXT. D&C VIOLET NO. 2 (UNII: G5UX3K0728)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60193-202-01	236 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product		
2	NDC:60193-202-03	1000 mL in 1 CARTRIDGE; Type 0: Not a Combination Product		
3	NDC:60193-202-05	1800 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product		
4	NDC:60193-202-07	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	05/01/2014	

Labeler - Pacific World Corporation (089693097)

Revised: 10/2014 Pacific World Corporation