8.75OZ ANTIBACTERIAL FOAM HANDSOAP - OCEAN BREEZE- benzalkonium chloride gel
Hero Brands, 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.

90114-253 259 ml Zorin Pharmaceutical Technology (Hangzhou) Co Ltd. 8.75oz ANTIBACTERIAL FOAM HAND SOAP - OCEAN BREEZE

Active Ingredient(s)

Benzalkonium Chloride 0.13% , Antibacterial

Purpose

Antibacterial, Hand SOAP

Use

For handwashing to decrease bacteria on skin.

Warnings

For external use only. Keep away from fire or flame

Do not use

- in children less than 2 months of age
- on open skin wounds

When using this product keep out of eyes. If contact with eyes occurs, rinse thoroughly with water.

Stop use and ask a doctor if irritation or rash develops.

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

Directions

- Wet hands
- Apply foaming soap to hands and work into rich lather
- Rinse hands thoroughly

Inactive ingredients

Disodium Cocoamphodiacetate,Water,Glycerin,Citric Acid, EDTA-2Na,CAB-35,Fragrance

Package Label - Principal Display Panel

宽43.606mm 高105.727mm 宽88.511 mm 高105.692mm

8.75OZ ANTIBACTERIAL FOAM HANDSOAP - OCEAN BREEZE

benzalkonium chloride gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:90114-253
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.013 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
SHEA BUTTER (UNII: K49155WL9Y)	
EDETIC ACID (UNII: 9G34HU7RV0)	

WATER (UNII: 059QF0KO0R)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
GLYCERIN (UNII: PDC6A3C0OX)	
DISODIUM COCOAMPHODIACETATE (UNII: 18L9G3U51M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:90114-253-01	259 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/05/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	11/05/2020	

Labeler - Hero Brands, Inc. (117527162)

Registrant - Hero Brands, Inc. (117527162)

Establishment

Name	Address	ID/FEI	Business Operations
Zorin Pharmaceutical Technology (Hangzhou) Co Ltd.		554529819	manufacture(90114-253)

Revised: 12/2020 Hero Brands, Inc.