

**MEDPRIDE ANTIBACTERIAL HAND WASH- benzalkonium chloride liquid
Shield Line LLC**

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.

Antibacterial Hand Soap- Tropical Beach

Drug Facts

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Active Ingredient

Benzalkonium Chloride 0.13%(w/w)

Purpose

Antibacterial

Uses

- For washing hand 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

irritation and redness develops

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

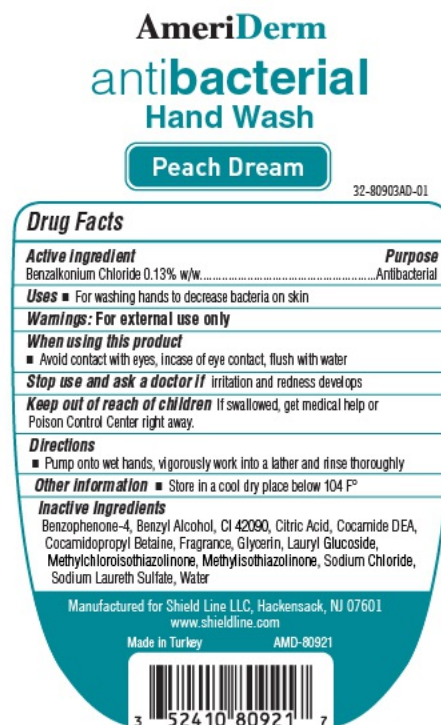
pump onto wet hands, vigorously work into lather rinse thoroughly

other information

Store in a cool dry place below 104 F

Inactive Ingredients

Benzophenone-4, Benzyl Alcohol, CI 42090, Citric Acid, Cocamide DEA, Cocamidopropyl Betaine, Fragrance, Glycerin, Lauryl Glucoside, Methylchloroisothiazolinone, Methylisothiazolinone, Sodium Chloride, Sodium Laureth Sulfate, Water



MEDPRIDE ANTIBACTERIAL HAND WASH			
benzalkonium chloride liquid			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:52410-8092
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)		BENZALKONIUM CHLORIDE	0.13 g in 100 g
Inactive Ingredients			
Ingredient Name			Strength
COCO DIETHANOLAMIDE (UNII: 92005F972D)			
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)			

SULISOBENZONE (UNII: 1W6L629B4K)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
LAURYL GLUCOSIDE (UNII: 76LN7P7UCU)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
GLYCERIN (UNII: PDC6A3C0OX)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
WATER (UNII: 059QF0K00R)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52410-8092-1	400 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/04/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/04/2020	

Labeler - Shield Line LLC (078518916)

Revised: 1/2022

Shield Line LLC