BETADINE FOAM CLEANSER- benzalkonium chloride aerosol, foam
Atlantis Consumer Healthcare, Inc.

BETADINE®
HAND & SKIN CLEANSER

Drug Facts

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antiseptic

Uses

- antimicrobial skin cleanser to help reduce bacteria that can potentially cause disease
- helps to prevent the spread of bacteria

Warnings

For external use only

When using this product

do not use in or near eyes.

Stop use and ask a doctor if

irritation and redness develop.

Keep out of reach of children.

If swallowed, seek medical attention or contact a Poison Control Center immediately.

Directions

- wet area to be cleaned with clean, running water
- apply appropriate amount of product
- lather and scrub for at least 1 seconds
- rinse thoroughly with running water
- dry completely

Other information

Store in a cool, dry place below 104° F (40° C)

Inactive ingredients

Water, glycerin, cocamidopropyl betaine, lauramine oxide, ethylene glycol phenyl ether, fragrance, cocamide MEA, tetrasodium EDTA, PEG-150 distearate, aloe barbadensis leaf juice.

Questions?

1-833-288-2684

Package Labels

BETADINE®
HAND & SKIN
CLEANSER

Antimicrobial
Foam Skin Cleanser

Hospital Grade,
Broad Spectrum
Germ Killing*

Great for Hand
& Skin Hygiene

Skin Softening
Formula
with Aloe

HAND WASHING

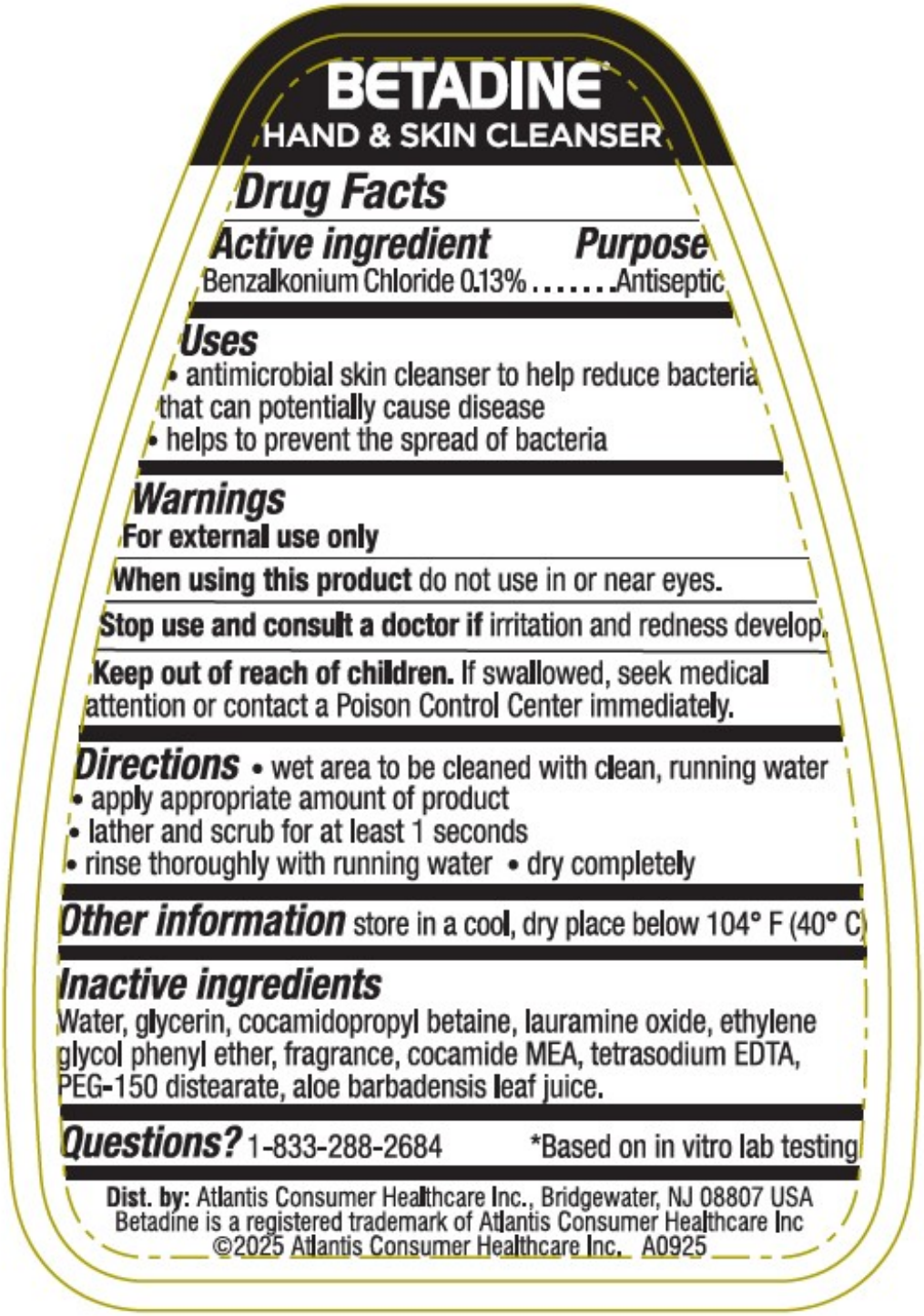
GENERAL SKIN CLEANSING

8 fl oz (237 mL)

BETADINE®
HAND & SKIN CLEANSER

*Based on in vitro lab testing

Dist. by: Atlantis Consumer Healthcare Inc., Bridgewater, NJ 08807 USA
Betadine is a registered trademark of Atlantis Consumer Healthcare Inc
 A0925



BETADINE®
HAND & SKIN
CLEANSER

Antimicrobial Foam
Skin Cleanser

NEW!

Hospital Grade,

Broad Spectrum
Germ Killing*

Kills 99.9% of 20
Different Bacteria
in Seconds*

Great for Hand
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BETADINE FOAM CLEANSER

benzalkonium chloride aerosol, foam

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67618-184
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 mL

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
Glycerin (UNII: PDC6A3C0OX)	
Cocamidopropyl Betaine (UNII: 5OCF3O11KX)	
Lauramine Oxide (UNII: 4F6FC4MI8W)	
Phenoxyethanol (UNII: HIE492ZZ3T)	
Coco Monoethanolamide (UNII: C80684146D)	
Edetate Sodium (UNII: MP1J8420LU)	
Peg-150 Distearate (UNII: 6F36Q0I0AC)	
Aloe Vera Leaf Juice (UNII: RUE8E6T4NB)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67618-184-08	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/01/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	02/01/2026	

Labeler - Atlantis Consumer Healthcare, Inc. (118983925)

Revised: 12/2025 Atlantis Consumer Healthcare, Inc.