MERINGUE FOAMING ANTIBACTERIAL HAND- benzalkonium chloride solution
United Laboratories, 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.

Drug Facts

Active ingredient

BENZALKONIUM CHLORIDE, 0.13%

Purpose

Antibacterial

Uses

for handwashing to reduce 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 develop and persist for more than 3 days.

Keep out of reach of children.

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

Directions

apply foaming cleanser to dry hands
rub hands together to spread lather
wash for 15-20 seconds
rinse & dry hands thoroughly

Inactive ingredients

AQUA (WATER), GLYCERIN, LAURAMINE OXIDE, BUTYLENE GLYCOL, LACTIC ACID, SALICYLIC ACID, PARFUM (FRAGRANCE), GREEN 5 (CI 61570), YELLOW 5 (CI 19140).

UNITED LABORATORIES

United 758
MERINGUE
Foaming Antibacterial Hand Soap
PRECAUCION AL USARIO: Si usted no pede leer Ingles, pregunte a alguien que le traduzca esta etigueta para usted antes de uso.
A high-quality, foam soap that is 99.99% effective against common bacteria.
An SDS for this product is available through United's website, www.unitedlabsinc.com, providing 24 hour access. Please read the SDS carefully and follow all directions when using or handling this product. Never reuse empty containers. Incompatible materials may adversely react.
NDC 63998-010-27
Manufactured in U.S.A.

1017

1 L (33.8 fl oz)

Sold By: UNITED LABORATORIES, INC.

320 37th Avenue ● St. Charles, IL 60174 ● 1-800-323-2594 ● www.unitedlabsinc.com

MERINGUE FOAMING ANTIBACTERIAL HAND

benzalkonium chloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63998-010
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)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
LACTIC ACID (UNII: 33X04XA5AT)	
SALICYLIC ACID (UNII: O414PZ4LPZ)	
D&C GREEN NO. 5 (UNII: 8J6RDU8L9X)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:63998-010-27	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/11/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	01/11/2018	

Labeler - United Laboratories, Inc. (001759737)