PIONEER ECLIPSE TOUCHE FOAMING NON-ALCOHOL HAND SANITIZERbenzalkonium chloride solution CWGC LA 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.

CWGC - Pioneer Eclipse Touché Foaming Non-Alcohol Hand Sanitizer (70415-305)

Active ingredient

BENZALKONIUM CHLORIDE 0.13%

Purpose

Antibacterial

Uses

• For handwashing to reduce bacteria on the skin. Recommended for repeated use.

Warnings

For external use only

Avoid contact with eyes. In case of eye contact, flush eyes with water.

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

Stop use and ask a doctor if irritation or redness develops and persists.

Directions

- Apply foam sanitizer to hands.
- Rub over surfaces of both hands for 15 seconds.
- No rinsing required.

Inactive ingredients

WATER, COCO-GLUCOSIDE, LAURTRIMONIUM CHLORIDE, COCAMIDOPROPYLAMINE OXIDE, CITRIC ACID, FRAGRANCE.





Touché TOAMING NON-ALCOHOL HAND SANITIZER

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Sparta, NC 28675 • USA Fax: 1-336-372-2895

Corp.

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Amano Pioneer Eclipse® Corp

Drug Facts

Active IngredientPurpose

Benzalkonium Chloride 0.13% Antimicrobial

Uses For handwashing to decrease bacteria on the skin. Recommended for repeat use.

Warnings

For external use only.

Avoid contact with eyes - In case of eye contact, flush eyes with water.

Keep out of reach of children. If swallowed, get immediate medical attention. Stop use and ask doctor if irritation or redness develops and persists.

Directions Apply foam sanitizer to hands. Rub over surfaces of both hands for 15 seconds. No rinsing required.

Inactive ingredients Water, Coco-Glucoside, Laurtrimonium Chloride, Cocamidopropylamine Oxide, Citric Acid, Fragrance.





CONTENTS: 1 Liter (33.8 fl. oz.)





PIONEER ECLIPSE TOUCHE FOAMING NON-ALCOHOL HAND SANITIZER

benzalkonium chloride solution

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Product Type HUMAN OTC DRUG Item Code (Source) NDC:70415-305

TOPICAL Route of Administration

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
	BENZALKONIUM CHLORIDE	0.13 g in 100 mL

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
COCO GLUCOSIDE (UNII: ICS790225B)		
LAURTRIMONIUM CHLORIDE (UNII: A81MSI0FIC)		
COCAMIDOPROPYLAMINE OXIDE (UNII: M4SL82J7HK)		
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)		

ı	Packaging				
	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
	1 NDC:70415-305-01	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/04/2017		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	08/04/2017		

Labeler - CWGC LA Inc. (034967904)

Revised: 9/2023 CWGC LA Inc.