

**PIONEER ECLIPSE TOUCHE FOAMING HAND SANITIZER- ethyl alcohol liquid
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.

Pioneer Eclipse Touché Foaming Hand Sanitizer (70415-302)

Active ingredient

Ethyl Alcohol 70 %

PURPOSE

ANTIBACTERIAL

Uses

- For sanitizing to reduce bacteria on the skin.

Warnings

For external use only

Flammable :Keep away from fire or flame

When using this product avoid contact with eyes. In case of eye contact, flush with water

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

Directions

- Apply one pump of foaming cleanser to dry hands.
- Rub into skin.
- No rinsing required.

Inactive ingredients

Water, Isopropyl Alcohol, Propylene Glycol, PEG-12 Dimethicone Crosspolymer, BIS-PEG-12 Dimethicone.

KILLS 99% OF GERMS INSTANTLY

ITEM #3079L4 · HAND SANITIZER



Touche™ FOAMING HAND SANITIZER

1 Eclipse Road, PO Box 909
Sparta, NC 28675 • USA
Fax: 1-336-372-2895

Amano Pioneer Eclipse® Corp.
1-800-367-3550 • 1-336-372-8080
www.pioneerclipse.com

© 2017 Amano Pioneer Eclipse Corp. 3079L4-0617

Drug Facts

Active Ingredient	Purpose
Ethyl Alcohol 70%.....	Antibacterial

Uses ■ For sanitizing to reduce bacteria on the skin.

Warnings
For external use only.

Flammable: Keep away from fire or flame.

When using this product, avoid contact with eyes. In case of eye contact, flush with water.

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

Directions
 ■ Apply one pump of foaming cleanser to dry hands. ■ Rub into skin.
 ■ No rinsing required.

Inactive ingredients Water, Isopropyl Alcohol, Propylene Glycol, PEG-12 Dimethicone Crosspolymer, Bis-PEG-12 Dimethicone.



CONTENTS:
1 Liter (33.8 fl. oz.)

HAND SANITIZER



PIONEER ECLIPSE TOUCHE FOAMING HAND SANITIZER

ethyl alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70415-302
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PEG-12 DIMETHICONE (300 CST) (UNII: ZEL54N6W95)	
BIS-PEG-12 DIMETHICONE (70 MPA.S) (UNII: 2JDK5W22H4)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70415-302-01	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/01/2016	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	02/01/2016	

Labeler - CWGC LA Inc. (034967904)

Revised: 9/2023

CWGC LA Inc.