EVERSOFT FOAMING HAND SANITIZER- ethyl alcohol liquid
Teh Tung Corporation

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.

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eversoft Foaming Hand Sanitizer

Drug Facts

Active Ingredient

Ethyl Alcohol, 73%

Purpose

Antimicrobial

Use

• Hand sanitizing to help reduce bacteria on the skin • Recommended for repeated use

Directions

Apply liberally to the hands and gently rub until dry.

Other Information

• Store above 0°C (32°F)

Inactive Ingredients

Purified Water, Dimethicone, Isopropyl Alcohol

Warnings

Flammable, keep away from fire or flame.

For external use only.

When using this product avoid contact with the eyes. In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or redness appears and lasts.

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

Foaming Hand Sanitizer - Alcohol Form

*Patented formula effectively eliminates 99.99% of tested germs, bacteria and viruses in as little as 15 seconds; long-term protection for up to 12 hrs.

Distributed by:

Draco Hygienic Products Inc.

716 S. Bon View Ave, Ontario, CA 91761

www.draco.com

Emergency: Chemtel 800-2553924

WARNING: FLAMMABLE LIQUID AND VAPOR.

CAUSES SERIOUS EYE IRRITATION. HARMFUL IF SWALLOWED.

Packaging

EVERSOFT FOAMING HAND SANITIZER
ethyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:78168-003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	73 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:78168-003-55	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/01/2020	
2	NDC:78168-003-56	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	11/01/2020	

Labeler - Teh Tung Corporation (023729484)

Revised: 10/2020 Teh Tung Corporation