KLEENLINE- 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.

KLEENLiNE™ FOAM HAND SANITIZER

Drug Facts

Active ingredient

Ethyl Alcohol 73%

Purpose

Antimicrobial

Uses

• 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

For external use only.

Flammable, keep away from fire or flame.

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.

Questions? Call **1-800-995-4466**

BY **WAXIE**

FOAM HAND SANITIZER

ETHANOL ALCOHOL

INSTANTLY KILLS 99.99% OF TESTED GERMS

FRAGRANCE-FREE

PROUDLY MADE IN USA

Distributed by:

WAXIE Sanitary Supply
San Diego, CA 92123

*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.

Packaging

KLEENLINE

ethyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:78168-628
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:78168-628-88	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/15/2020	
2	NDC:78168-628-99	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/15/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/15/2020	

Labeler - Teh Tung Corporation (023729484)

Revised: 10/2020 Teh Tung Corporation