MOREX- 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.

***morex*® 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.

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.

Other information

Store above 0°C (32°F)

Inactive ingredients

Purified water, Ethyl alcohol

Questions? Call **1-562-903-8000**

ALCOHOL-BASED

INSTANTLY KILLS 99.99% OF TESTED GERMS

FRAGRANCE -FREE

PROUDLY MADE IN USA

Distributed by:

TEH TUNG CORPORATION

Santa Fe Springs, CA 90670

*Patents formula effectively eliminates 99.99% of tested germs, bacteria and viruses in as little as 15 seconds

Packaging

MOREX

ethyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:78168-868
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)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:78168-868-01	55 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/15/2020	
2	NDC:78168-868-02	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/15/2020	
3	NDC:78168-868-03	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/15/2020	
4	NDC:78168-868-04	120 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/15/2020	
5	NDC:78168-868-07	200 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/15/2020	
6	NDC:78168-868-08	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/15/2020	
7	NDC:78168-868-19	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/15/2020	
8	NDC:78168-868-24	710 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/15/2020	
9	NDC:78168-868-32	950 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/15/2020	
10	NDC:78168-868-69	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/15/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/15/2020	

Labeler - Teh Tung Corporation (023729484)

Revised: 6/2020 Teh Tung Corporation