MOREX HAND SANITIZER- benzalkonium chloride solution
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

Benzalkonium Chloride 0.13%

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.

Inactive ingredients

Purified water, D-glucopyranose, Oligomeric, Decyl octyl glycosides, Glycerol, Methylisothiazolinone, and Citric acid

Warnings

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.

Other information

Store above 0°C (32°F)

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

ALCOHOL-FREE

INSTANTLY KILLS 99.99% OF TESTED GERMS

HYPOALLERGENIC

PROUDLY MADE IN USA

Distributed by:

TEH TUNG CORPORATION

Santa Fe Springs, CA 90670

*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

MOREX HAND SANITIZER

benzalkonium chloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:78168-608
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
2,3 DI-O-METHYL-D-GLUCOSE (UNII: 3F1Y194PDX)	
CAPRYL OLIGOGLUCOSIDE (UNII: VXR09E583M)	
CAPRYLYL/CAPRYL OLIGOGLUCOSIDE (UNII: E00JL9G9K0)	
GLYCERIN (UNII: PDC6A3C0OX)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:78168-608-01	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/15/2020	
2	NDC:78168-608-02	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/15/2020	
3	NDC:78168-608-04	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/15/2020	
4	NDC:78168-608-08	120 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/15/2020	
5	NDC:78168-608-09	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/15/2020	
6	NDC:78168-608-12	350 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/15/2020	
7	NDC:78168-608-16	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/15/2020	
8	NDC:78168-608-24	710 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/15/2020	
9	NDC:78168-608-32	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/15/2020	
10	NDC:78168-608-89	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