HAND SANITIZER- hand sanitizer liquid
Guangdong Boxi High-tech Co.,Ltd.

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.

HAND SANITIZER

Active Ingredient(s)

Alcohol 62.0%

Purpose

Antimicrobial

Uses

Hand Sanitizer to help reduce bacteria on skin

WARNINGS

For external use only. Flammable. Keep away from heat or flame

When using this product

When using this product, do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water Stop use and ask doctor if irritation or rash appears and lasts. Keep out of reach for children if swallowed, get medical help or contact a Poison Control Center right away

Directions

- Children under 6 years of age should be supervised when using this product
- Place enough Product in your palm to thoroughly spread on both hands and bub into the skin until dry

Other information

Store below 30 C(86 F)

Inactive ingredients

Water(Aqua), Acrylates copolymer, Triethanolamine, Fragrance,Aloe barbadensis leaf extract, Tocopherol, Denatonium benzoate.

Package Label - Principal Display Panel

HAND SANITIZER

hand sanitizer liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75702-006
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
WATER (UNII: 059QF0KO0R)	
ALOE (UNII: V5VD430YW9)	
TROLAMINE (UNII: 9O3K93S3TK)	
BUTYL ACRYLATE/METHYL METHACRYLATE/METHACRYLIC ACID COPOLYMER (18000 MW) (UNII: JZ1374NL9E)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75702-006-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/22/2020	
2	NDC:75702-006-02	40 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/22/2020	
3	NDC:75702-006-03	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/22/2020	
4	NDC:75702-006-04	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/22/2020	
5	NDC:75702-006-05	80 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/22/2020	
6	NDC:75702-006-06	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/22/2020	
7	NDC:75702-006-07	120 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/22/2020	
8	NDC:75702-006-08	200 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/22/2020	
9	NDC:75702-006-09	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/22/2020	
10	NDC:75702-006-10	280 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/22/2020	
11	NDC:75702-006-11	300 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/22/2020	
12	NDC:75702-006-12	420 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/22/2020	
13	NDC:75702-006-13	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/22/2020	
14	NDC:75702-006-14	600 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/22/2020	
15	NDC:75702-006-15	800 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/22/2020	
16	NDC:75702-006-16	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/22/2020	
17	NDC:75702-006-17	2000 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/22/2020	
18	NDC:75702-006-18	3000 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/22/2020	
19	NDC:75702-006-19	4000 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/22/2020	
20	NDC:75702-006-20	5000 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/22/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/22/2020	

Labeler - Guangdong Boxi High-tech Co.,Ltd. (542874709)

Establishment

Name	Address	ID/FEI	Business Operations
Guangdong Boxi High-tech Co.,Ltd.		542874709	manufacture(75702-006)

Revised: 4/2020 Guangdong Boxi High-tech Co.,Ltd.