HAND SANITIZER- alcohol gel
NingBo Huize Commodity 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

Ethyl Alcohol 62%

Purpose

Antiseptic

Use

Decreases bacteria on hands

Warning

for external use only

Do not use on broken skin & around eyes

When using this product
■Keep away from fire or flame
■Using other topical acne drugs at the same time or right after use of this product may increase dryness or irritation of the skin . Only one drug should be used unless directed by a doctor.

Keep out of reach of children

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

Directions

Squeeze onto hands and rub hands and rub hands together briskly until dry

INACTIVE INGREDIENT

Deionized Water, glycerin , Propylene Glycol, aloe vera gel,fragrance,DMDMH

Drug Facts	Drug Facts(continued)
Active Ingredient **Purpose** Ethyl Alcohol 62% ..... Antiseptic **Uses** ■Decreases bacteria on hands **Warnings** For external use only	**Directions** ■Squeeze onto hands and rub hands and rub hands together briskly until dry
Do not use ■on broken skin & around eyes	**Other information:** store at 20-25 ℃（68-77 ℉）
When using this product ■Keep away from fire or flame ■Using other topical acne drugs at the same time or right after use of this product may increase dryness or irritation of the skin . Only one drug should be used unless directed by a doctor.	**Inactive ingredients** Deionized Water, glycerin , Propylene Glycol, aloe vera gel,fragrance,DMDMH
Stop use and ask a doctor if rash occurs	
Keep out of reach of children .If swallowed ,get medical help or contact a Poison on Control Center immediately	

HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71011-003
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
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PROPYLENE GLYCOL DIACETATE (UNII: 5Z492UNF9O)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71011-003-01	5 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/20/2016	
2	NDC:71011-003-02	8 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/20/2016	
3	NDC:71011-003-03	10 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/20/2016	

4	NDC:71011-003-04	15 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/20/2016	
5	NDC:71011-003-05	20 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/19/2016	
6	NDC:71011-003-06	30 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/19/2016	
7	NDC:71011-003-07	50 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/20/2016	
8	NDC:71011-003-08	60 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/20/2016	
9	NDC:71011-003-09	80 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/20/2016	
10	NDC:71011-003-10	100 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/20/2016	
11	NDC:71011-003-11	150 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/20/2016	
12	NDC:71011-003-12	200 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/20/2016	
13	NDC:71011-003-13	250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/20/2016	
14	NDC:71011-003-14	300 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/20/2016	
15	NDC:71011-003-15	400 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/20/2016	
16	NDC:71011-003-16	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/20/2016	
17	NDC:71011-003-17	600 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/20/2016	
18	NDC:71011-003-18	700 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/20/2016	
19	NDC:71011-003-19	800 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/20/2016	
20	NDC:71011-003-20	900 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/20/2016	
21	NDC:71011-003-21	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/20/2016	
22	NDC:71011-003-22	2000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/20/2016	
23	NDC:71011-003-23	5 mL in 1 POUCH; Type 0: Not a Combination Product	10/20/2016	
24	NDC:71011-003-24	8 mL in 1 POUCH; Type 0: Not a Combination Product	10/20/2016	
25	NDC:71011-003-25	25 mL in 1 POUCH; Type 0: Not a Combination Product	10/20/2016	
26	NDC:71011-003-26	15 mL in 1 POUCH; Type 0: Not a Combination Product	10/20/2016	
27	NDC:71011-003-27	20 mL in 1 POUCH; Type 0: Not a Combination Product	10/20/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/09/2016	

Labeler - NingBo Huize Commodity Co.,Ltd. (544434795)

Establishment

Name	Address	ID/FEI	Business Operations
NingBo Huize Commodity Co.,Ltd.		544434795	manufacture(71011-003)

Revised: 1/2018

NingBo Huize Commodity Co.,Ltd.