

HAND SANITIZER- ethyl alcohol gel
Hangzhou Huiji Biotechnology 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.

73788-004 75% alcohol hand sanitizer gel

Active Ingredient

Ethyl Alcohol 75%

Purpose

disinfectant

Use

■ Hand sanitizer to help decrease bacterial on
the skin

■ When water, soap & towel are not
available.

■ Recommended for repeated use,

WARNINGS

For external use only ■ Highly inflammable
Keep away from heat, sparks, open flame, hot surfaces.
Do not apply around eyes. Do not use in ears & mouth.
Avoid contact with eyes
If in eyes: rinse well with clean water
Stop use and seek medical advice or attention if
irritation persists

keep out of reach of children

Keep out of reach of children. Children must be
supervised in use of this product

Directions

■ Squeeze the bottle after removing the lid and thoroughly
spread on both hands. Rub into skin until dry

Other Information Avoid contact with fabrics as could cause discoloration

Inactive ingredients

Water, Glycerin, Aloe barbadensis leaf extract, Carbomer. Tocopheryl Acetate, Triethanolamine



HAND SANITIZER			
ethyl alcohol gel			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC: 73788-004
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)		ALCOHOL	75 mL in 100 mL	
Inactive Ingredients				
Ingredient Name			Strength	
WATER (UNII: 059QF0KO0R)				
GLYCERIN (UNII: PDC6A3C0OX)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
TROLAMINE (UNII: 9O3K93S3TK)				
CARBOMER 934 (UNII: Z135WT9208)				
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73788-004-01	200 in 1 CARTON	05/29/2020	
1		30 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:73788-004-02	150 in 1 CARTON	05/29/2020	
2		60 mL in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:73788-004-03	100 in 1 CARTON	05/29/2020	
3		100 mL in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:73788-004-04	24 in 1 CARTON	05/29/2020	
4		237 mL in 1 BOTTLE; Type 0: Not a Combination Product		
5	NDC:73788-004-05	20 in 1 CARTON	05/29/2020	
5		500 mL in 1 BOTTLE; Type 0: Not a Combination Product		
6	NDC:73788-004-06	18 in 1 CARTON	05/29/2020	
6		1000 mL in 1 BOTTLE; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	05/29/2020		

Labeler - Hangzhou Huiji Biotechnology Co., Ltd. (526893497)

Establishment

Name	Address	ID/FEI	Business Operations
Hangzhou Huiji Biotechnology Co., Ltd.		526893497	manufacture(73788-004)

Revised: 2/2022

Hangzhou Huiji Biotechnology Co., Ltd.