GRAPE FRUIT HAND SANITIZER- ethyl alcohol gel
SHANDISHI Biological Technology 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.

77753-005 Grape Fruit Hand Sanitizer 70% Alcohol

DRUG FACTS

Active ingredient

Ethyl Alcohol 70% v/v

Purpose

Antiseptic

Uses

For Hand Sanitizing

Warnings

Warnings: For external use only-hands.
Flammable: Keep away from heat and flame.
When using this product Keep out of eyes. In case of
contact with eyes, flush thoroughly with water.●Avoid contact with
broken skin.●Do not inhale or ingest.
Stop use and ask a doctor
if skin irritation develops.

Keep out of reach of children.

Keep out of reach of children

If swallowed, get medical help or call a Poison Control Center right away.

Directions

Wet hands thoroughly with product and allow to
dry without wiping●For children under 6 use only under adult
Supervision.●Not recommended for infants.

Other information

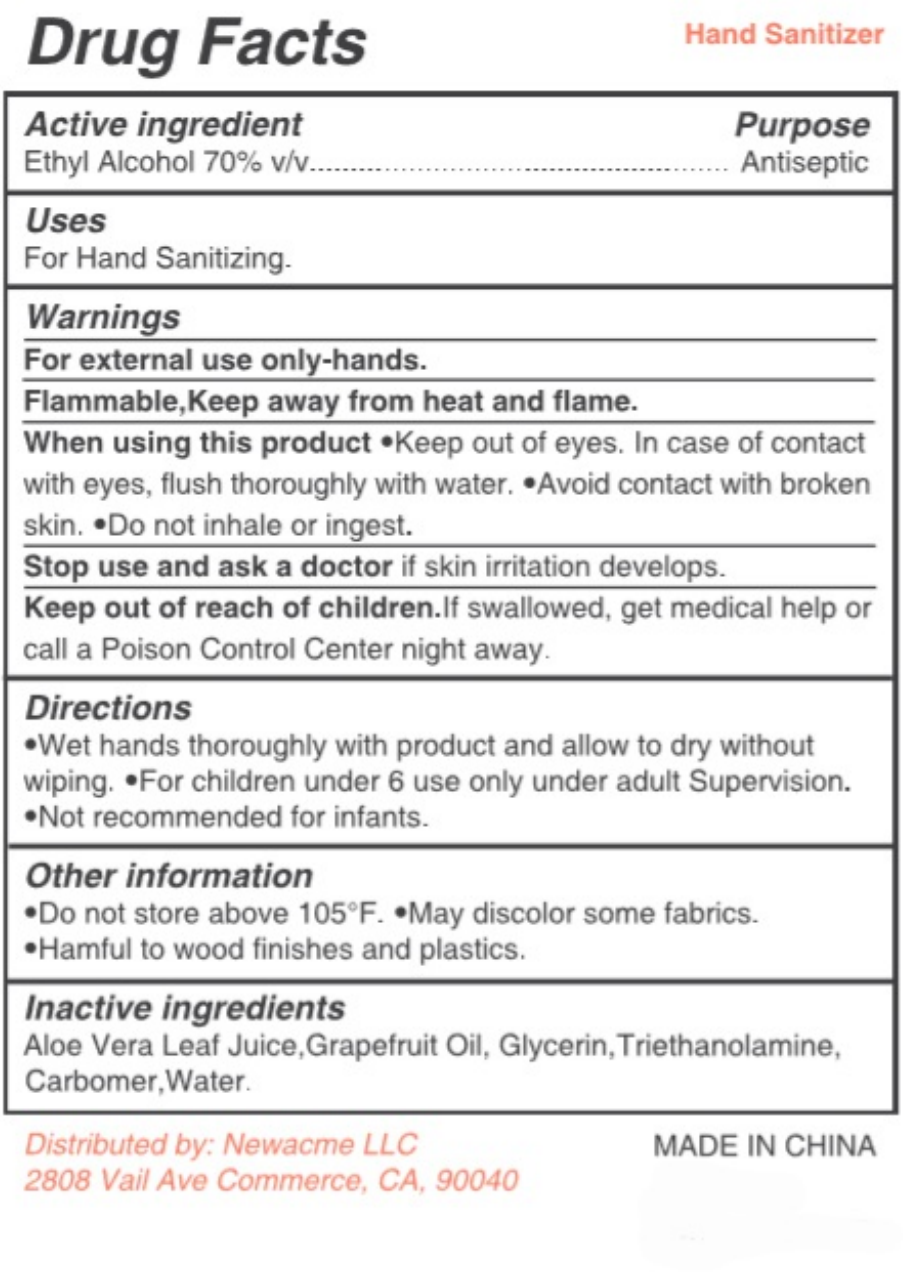
Do not store above 105°F
May discolor some fabrics

Hamful to wood finishes and plastics.

Inactive ingredients

Aloe Vera Leaf Juice,Grapefruit Oil, Glycerin,Triethanolamine,Carbomer,Water.

Package Labeling



Drug Facts

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Distributed by: Newacme LLC
2808 Vail Ave Commerce, CA, 90040

MADE IN CHINA

GRAPE FRUIT HAND SANITIZER

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:77753-005
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
VITIS VINIFERA FRUIT OIL (UNII: YQ5Q4Y2Z8U)	
GLYCERIN (UNII: PDC6A3C0OX)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77753-005-01	12 in 1 BOX	07/15/2020	
1		30 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:77753-005-02	12 in 1 BOX	07/15/2020	
2		53 mL in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:77753-005-03	12 in 1 BOX	07/15/2020	
3		60 mL in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:77753-005-04	2 in 1 BOX	07/15/2020	
4		250 mL in 1 BOTTLE; Type 0: Not a Combination Product		
5	NDC:77753-005-05	12 in 1 BAG	07/15/2020	
5		300 mL in 1 BOTTLE; Type 0: Not a Combination Product		
6	NDC:77753-005-06	8 in 1 BOX	07/15/2020	
6		350 mL in 1 BOTTLE; Type 0: Not a Combination Product		
7	NDC:77753-005-07	8 in 1 BOX	07/15/2020	
7		400 mL in 1 BOTTLE; Type 0: Not a Combination Product		
8	NDC:77753-005-08	8 in 1 BOX	07/15/2020	
8		500 mL in 1 BOTTLE; Type 0: Not a Combination Product		
9	NDC:77753-005-09	4 in 1 BOX	07/15/2020	
9		750 mL in 1 BOTTLE; Type 0: Not a Combination Product		
10	NDC:77753-005-10	4 in 1 BOX	07/15/2020	
10		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	07/15/2020	

Labeler - SHANDISHI Biological Technology Co., Ltd. (411868994)

Establishment

Name	Address	ID/FEI	Business Operations
SHANDISHI Biological Technology Co., Ltd.		411868994	manufacture(77753-005)

Revised: 7/2020

SHANDISHI Biological Technology Co., Ltd.