

HAND SANITIZER- hand sanitizer spray
Dongguan Mingyi Medical Products 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.

74046-004
HAND SANITIZER SPRAY

Active Ingredients

Ethyl Alcohol 75%

Purpose

Antimicrobial

Uses

Hand sanitizer to help reduce bacteria on skin.

Warnings

For external use only. Keep out of reach of children. Children must be supervised if using this product. Flammable. Keep away from fire or flame .Avoid contact with eyes. Discontinue use if irritation or redness occurs , If irritation continues for more than 72 hrs , please consult a doctor.

Directions

Spray enough product on hands cover all surfaces. Rub hands together until dry. Supervise children under 6 years of age when using this product to avoid swallowing.

Other Information

store below 105. 8°F (41°C)

may discolor certain fabrics or surfaces

Inactive Ingredients

Deionized water , Glycerol

Package Label - Principal Display Panel

Drug Facts

Active Ingredients

Ethyl alcohol 75% v/v,

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Manufacturer: Dongguan Mingyi medical supplies Co, Ltd

Address : Room 201, 1f, No. 187, Tianheng Road, Shahukou management zone, Changping Town, Dongguan City, Guangdong Province.

Made in China

99.9%
Effective bacteriostatic rate

**HAND
SANITIZER
SPRAY**

75% Alcohol



0.33FLoz /10ML

EXTERNAL USE

Drug Facts

Active Ingredients	PURPOSE
Ethyl alcohol 75% v/v,	Antimicrobial

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**HAND
SANITIZER
SPRAY**

75% Alcohol



0.52FLoz /15ML

EXTERNAL USE

Drug Facts

Active Ingredients

Ethyl alcohol 75% v/v,

PURPOSE

Antimicrobial

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**HAND
SANITIZER
SPRAY**

75% Alcohol



0.70FLoz /20ML

EXTERNAL USE

Drug Facts

Active Ingredients

Ethyl alcohol 75% v/v,

PURPOSE

Antimicrobial

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**HAND
SANITIZER
SPRAY**

75% Alcohol



1.05FLoz /30ML

EXTERNAL USE

Drug Facts

Active Ingredients

Ethyl alcohol 75% v/v,

PURPOSE

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**HAND
SANITIZER
SPRAY**

75% Alcohol



1.40FLoz /40ML

EXTERNAL USE

Drug Facts

Active Ingredients

Ethyl alcohol 75% v/v,

PURPOSE

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**HAND
SANITIZER
SPRAY**

75% Alcohol



1.75FLoz /50ML

EXTERNAL USE

Drug Facts

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Ethyl alcohol 75% v/v,

PURPOSE

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**HAND
SANITIZER
SPRAY**

75% Alcohol



2.11FLoz /60ML

EXTERNAL USE

Drug Facts

Active Ingredients

Ethyl alcohol 75% v/v,

PURPOSE

Antimicrobial

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**HAND
SANITIZER
SPRAY**
75% Alcohol



8.30FLoz /236ML

EXTERNAL USE

Drug Facts

Active Ingredients

Ethyl alcohol 75% v/v,

PURPOSE

Antimicrobial

Uses Hand sanitizer to help reduce bacteria on skin.

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**HAND
SANITIZER
SPRAY**

75% Alcohol



8.79FLoz /250ML

EXTERNAL USE

Drug Facts

Active Ingredients

Ethyl alcohol 75% v/v,

PURPOSE

Antimicrobial

Uses Hand sanitizer to help reduce bacteria on skin.

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**HAND
SANITIZER
SPRAY**
75% Alcohol



17.59FLoz / 500ML

EXTERNAL USE

HAND SANITIZER

hand sanitizer spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74046-004
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)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74046-004-01	10 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/15/2022	
2	NDC:74046-004-02	15 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/15/2022	
3	NDC:74046-004-03	20 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/15/2022	
4	NDC:74046-004-04	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/15/2022	
5	NDC:74046-004-05	40 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/15/2022	
6	NDC:74046-004-06	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/15/2022	
7	NDC:74046-004-07	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/15/2022	
8	NDC:74046-004-08	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/15/2022	
9	NDC:74046-004-09	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/15/2022	
10	NDC:74046-004-10	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/15/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/15/2022	

Labeler - Dongguan Mingyi Medical Products Co., Ltd. (554526907)

Registrant - Dongguan Mingyi Medical Products Co., Ltd. (554526907)

Establishment

Name	Address	ID/FEI	Business Operations
Dongguan Mingyi Medical Products Co., Ltd.		554526907	manufacture(74046-004)

Revised: 1/2022

Dongguan Mingyi Medical Products Co., Ltd.