

MAURICES I AM HAPPY HAND SANITIZER- hand sanitizer gel
Shanghai Sibe Cosmeic 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.

Active Ingredient(s)

Ethyl Alcohol 70%

Purpose

Antiseptic

Uses

- to decrease bacteria on the skin that could cause disease
- recommend for repeated use

Warnings

For external use only: hands

Flammable, keep away from fire or flame.

When using this product

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- 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 and ask a doctor if

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- • irritation and redness develop
- • condition persists for more than 72 hours

Keep out of the reach of children

Keep out of the reach of children. If swallowed, get medical help or contact 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

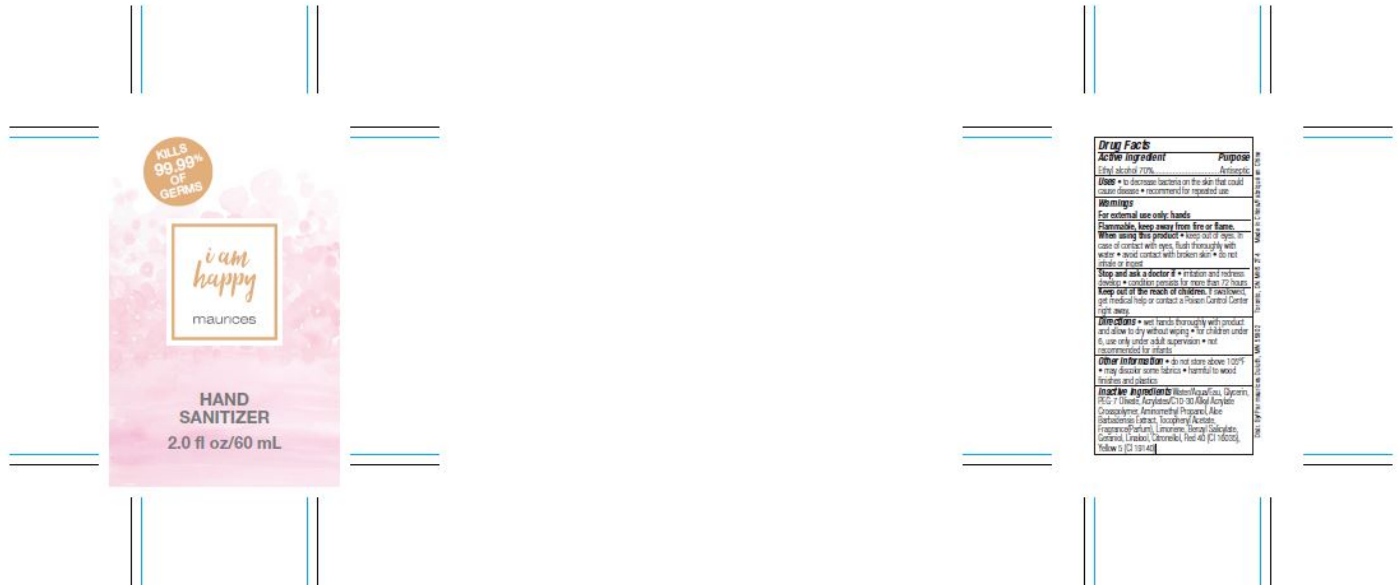
- do not store above 105°F
- may discolor some fabrics
- harmful to wood finishes and plastics

Inactive Ingredients

Water/Aqua/Eau, Glycerin, PEG-7 Oliviate, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aminomethyl Propanol, Aloe Barbadensis Extract, Tocopheryl Acetate, Fragrance(Parfum), Limonene, Benzyl Salicylate, Geraniol, Linalool, Citronellol, Red 40 (CI 16035), Yellow 5 (CI 19140)

Package Label/Principal Display Panel

NDC 79329-002-06; 60 ml



MAURICES I AM HAPPY HAND SANITIZER

hand sanitizer gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79329-002
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
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
GLYCERIN (UNII: PDC6A3C0OX)	
PEG-7 OLEATE (UNII: 7W4ZYQ7UVQ)	
BENZYL SALICYLATE (UNII: WAO5MKN9TU)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
GERANIOL (UNII: L837108USY)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	

WATER (UNII: 059QF0KO0R)	
.BETA.-CITRONELLOL, (R)- (UNII: P01OUT964K)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
ALOE VERA WHOLE (UNII: KIZ4X2EHYX)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79329-002-06	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/15/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/15/2020	

Labeler - Shanghai Sabei Cosmetic Co., Ltd. (542969967)

Establishment

Name	Address	ID/FEI	Business Operations
Shanghai Sabei Cosmetic Co., Ltd		542969967	manufacture(79329-002)

Revised: 7/2020

Shanghai Sabei Cosmetic Co., Ltd.