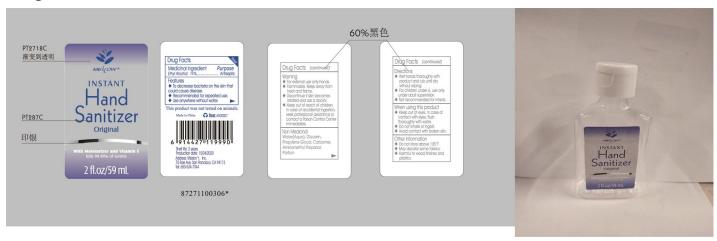
MELLOW 59ML HAND SANITIZER ORIGINAL- alcohol liquid Mission Y,Inc.

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.

Mellow 59ml hand sanitizer original

Drug Facts



Medicical Ingredient

Medicical Ingredient Purpose

Ethyl Alcohol 75% IV/VI Antiseptic

Features

To decrease bacteria on the skin that could cause disease.

Recommended for repeated use.

Use anywhere without water.

Recommended for repeated use.

use anywhere without water.

Warning

For external use only-hands.

Flammable, keep away from heat and flame.

Discontinue if skin becomes irritated and ask a doctor.

Keep out of reach of children. In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.

For external use only.

Flammable, keep away from heat and flame.

Discontinue if skin becomes irritated and ask a doctor.

Keep out of reach of children. In case of accidental ingestion, seek professional assistance or contact a poson control center immediately.

Non-Medicinal

Water, Glycerin, Propylene Glycol, Carbomer,, Aminomethyl propanol, Parfum.

Directions

Wet hands thoroughly with product and rub until dry without wiping

For children under 6, use only under adult supervision.

Not recommended for infants.

When using this product

keep out of eyes. In case of contact with eyes, flush thoroughly with water.

Do not inhale or ingest.

Avoid contact with broken skin.

Other information

Do not store above 105F.

May discolor some fabrics.

Harmful to wood finishes and plastics.

When using this product

keep out of eyes, In case of contact with eyes, flush thoroughly with water

Do not inhale or ingest

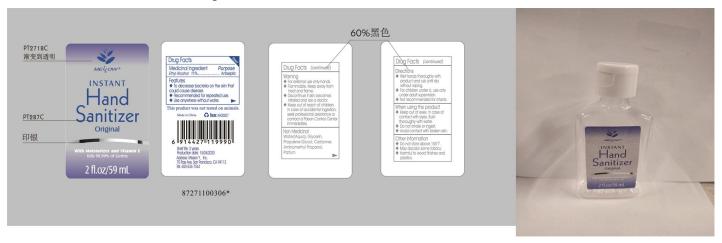
Avoid contact with broken skin.

other information

Do not store above 1050

May discolor some fabrics.

Harmful to wood finishes and plastics [



MELLOW 59ML HAND SANITIZER ORIGINAL

alcohol liquid

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75580-005	
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			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	0.01 mL in 100 mL			
LEMON (UNII: 24RS0A988O)	0.1 mL in 100 mL			
CARBOMER 940 (UNII: 4Q93RCW27E)	0.35 mL in 100 mL			
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	0.11 mL in 100 mL			
WATER (UNII: 059QF0KO0R)	28.42 mL in 100 mL			
GLYCERIN (UNII: PDC6A3C0OX)	0.01 mL in 100 mL			

Product Characteristics				
Color	white (transparant)	Score		
Shape		Size		
Flavor		Imprint Code		
Contains				

l	Packaging			
# Item Code Package Description		Marketing Start Date	Marketing End Date	
	1 NDC:75580-005- 01	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/20/2020	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	04/20/2020		

Labeler - Mission Y,Inc. (117492659)

Registrant - Mission Y,Inc. (117492659)

Establishment			
Name	Address	ID/FEI	Business Operations
Mission Y,Inc.		11749 26 59	label(75580-005)

Establishment				
Name	Address	ID/FEI	Business Operations	
Shenzhen Lantern Science Co.,Ltd.		421222423	manufacture(75580-005)	

Revised: 4/2020 Mission Y,Inc.