PASION INSTANT HAND SANITIZER- alcohol gel
MING FAI ENTERPRISE INTERNATIONAL 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.

Pasion Instant Hand Sanitizer

Drug Facts

Active Ingredient

Ethyl Alcohol 72%(v/v)

Purpose

Antiseptic

Uses

• to help reduce bacteria on the skin.

Warnings

- For external use only. • Flammable. • Keep away from fire or flame.

When using this product

• avoid contact with eyes. • If contact occurs, rinse thoroughly with water.

Stop using and ask a doctor if

• irritation or redness develops and lasts.

Keep out of reach of children

• In case of accidental ingestion, get medical help or contact a poison control center immediately.

Directions

• Squeeze a significant amount in your palm and rub hands until fully dry. • Rinse free.

Other information

• Store below 110°F (43°C).

Inactive ingredients

Water(Aqua), Glycerin, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Phenoxyethanol, Melaleuca Alternafolia (Tea Tree) Leaf Oil, Aminomethyl propanol.

Package Labeling:72135-031-50

Package Labeling:72135-031-30

Package Labeling:74548-031-10

PASION INSTANT HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:74548-031
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.72 mL in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
MELALEUCA ALTERNIFOLIA LEAF (UNII: G43C57162K)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:74548-031-50	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020	
2	NDC:74548-031-30	300 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020	
3	NDC:74548-031-10	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	05/01/2020	

Labeler - MING FAI ENTERPRISE INTERNATIONAL CO., LTD (667902568)

Revised: 4/2020 MING FAI ENTERPRISE INTERNATIONAL CO., LTD