### JIT HAND SANITIZERS- alcohol gel KMPHARMACEUTICAL 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.

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# **ACTIVE INGREDIENT**

Active ingredients: ALCOHOL 62.0%

# **INACTIVE INGREDIENT**

Inactive Ingredients:

Water, Glycerin, Butylene Glycol, Aloe Barbadensis Leaf Extract, Carbomer, Triethanolamine, Flavor

### PURPOSE

Purpose: SANITIZER

#### WARNINGS

Warnings:

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

When using this product do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.

Discontinue use if irritation or redness develops. If condition persists for more than 72 hours, consult a doctor.

# **KEEP OUT OF REACH OF CHILDREN**

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

#### Uses

Uses: Hand sanitizer to help reduce bacteria on the skin.

# Directions

Directions: Put enough product in your palm to cover hands and rub hands together briskly until dry. Children under 6 years of age should be supervised when using this product.

# PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



lcohol gel					
Product Inform	ation				
Product T ype		HUMAN OTC DRUG	em Code (So	urce) N	DC:50555-050
Route of Administ	ration	TOPICAL			
Active Ingredie	nt/Activ	e Moiety			
Ingredient Name				Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)				COHOL	37.2 g in 60 mL
Inactive Ingred	ients				
		Ingredient Name			Strength
Water (UNII: 059QF	)KO0R)	<u> </u>			
Glycerin (UNII: PDC	6A3C0OX	)			
Butylene Glycol (UI	MI: 3XUS8	5K0RA)			
ALOE VERA LEAF (	UNII: ZY8	1Z83H0X)			
CARBO MER HO MO	POLYME	R, UNSPECIFIED TYPE (UNII: 0A5MM307	FC)		
TROLAMINE (UNII:	9O3K93S	3TK)			
Packaging					
# Item Code		Package Description	Ma	nrketing Start Date	Marketing End Dat
$1 \begin{array}{c} NDC: 50555-050-\\ 02 \end{array}$	1 in 1 CA	RTON	03/01	/2020	
NDC:50555-050- 01 60 mL in 1 CONTAINER; Type 0: Not a Combination Product			1		
	forma	tion			
Marketing In			tion Mark	ating Start Data	Marketing End Date
Marketing In Marketing Categ	ory A	pplication Number or Monograph Cita		eting Start Date	Marketing End Date

Labeler - KMPHARMACEUTICAL Co.,Ltd. (688679158)

# Registrant - KMPHARMACEUTICAL Co.,Ltd. (688679158)

# Establishment

Name	Address	ID/FEI	<b>Business Operations</b>
KMPHARMACEUTICAL Co.,Ltd.		688679158	manufacture(50555-050)