LECHAT- hand sanitizer gel gel
Mega Creation 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.

LECHAT Sanitizer Gel

Active Ingredient(s)

Ethyl Alcohol 65% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer

Use

Hand Sanitizer to help reduce bacteria.

Warnings

For external use only.

Flammable.

Keep away from heat or flame

Do not use

- in children less than 2 months of age
- on open skin wounds

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.
Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

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

Directions

- Wet hands thoroughly with the product and rub it on the skin untill dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

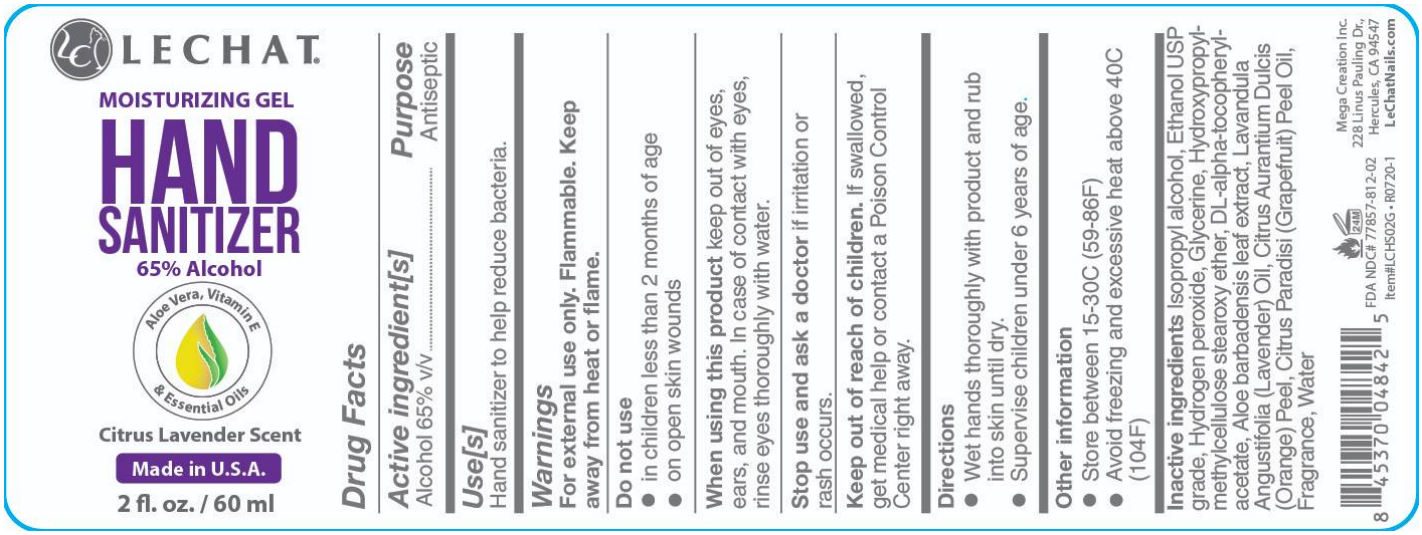
Other information

- Store between 15-30 C (59-86F)
- Avoid freezing & excessive heat above 40C (104F)

Inactive ingredients

glycerin, hydrogen peroxide, purified water, isopropyl alcohol, hydroxypropyl methylcellulose, dl alpha-tocopheryl acetate, Aloe barbadensis leaf extract, Lavandula Angustifolia oil, citrus Aurantium Dulcis peel, citrus paradisi peel oil, Fragrance

Package Label - Principal Display Panel



60mL NDC: 77857-812-02

LECHAT

hand sanitizer gel gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:77857-812
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	65 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**

GLYCERIN (UNII: PDC6A3C0OX)	
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
CITRUS PARADISI FRUIT OIL (UNII: 6A7N43E0OJ)	
LAVENDER OIL (UNII: ZBP1YXW0H8)	
ORANGE PEEL (UNII: TI9T76XD44)	
FRAGRANCE GREEN APPLE ORC2001072 (UNII: U9GH30P956)	
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77857-812-01	51 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/22/2020	
2	NDC:77857-812-02	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/22/2020	
3	NDC:77857-812-04	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/22/2020	
4	NDC:77857-812-08	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/22/2020	
5	NDC:77857-812-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/22/2020	
6	NDC:77857-812-32	946 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/22/2020	
7	NDC:77857-812-28	3780 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/22/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/22/2020	

Labeler - Mega Creation Inc. (014096208)

Registrant - Mega Creation Inc. (014096208)

Establishment

Name	Address	ID/FEI	Business Operations
Mega Creation Inc.		014096208	manufacture(77857-812)

Revised: 10/2022 Mega Creation Inc.