

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 New 60% Ethanol

Active Ingredient(s)

Ethyl Alcohol 60% 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.

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Directions

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

Other information

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

Inactive ingredients

Purified Water, IsopiopylAlcohol, Hydrogen Peroxide, Glycerin, Isopropyl Myristate, Acrylates/C10-30 Ally Acrylate Crosspolymer, Aminomethyl Propanol, Cocoyl Caprylocaprata, Aloe Barbadensis Leaf Extract, Tocopheryl-acetate, fragrance, Lavender Oil, Orange Oil, Grapefruit Oil

Package Label - Principal Display Panel

LECHAT.
MOISTURIZING GEL
HAND SANITIZER
60% Alcohol
Aloe Vera, Vitamin E & Essential Oils
Citrus Lavender Scent
MADE IN U.S.A.
1.75 fl. oz. / 51 ml

Drug Facts
Active ingredient[s] Purpose
Alcohol 60% v/v.....Antiseptic

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Item#LCHS01G-2
Mega Creation Inc.
FDA NDC# 228 Linus Pauling Dr.
Hercules, CA 94547
77857-832-01
R0621-1
LeChatMails.com

51 mL NDC: 77857-832-01

LECHAT			
hand sanitizer gel gel			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77857-832
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	60 mL in 100 mL	
Inactive Ingredients			
Ingredient Name	Strength		
GLYCERIN (UNII: PDC6A3C0OX)			
HYDROGEN PEROXIDE (UNII: BBX060AN9V)			
WATER (UNII: 059QF0KO0R)			
ALOE VERA LEAF (UNII: ZY81Z83H0X)			

LAVENDER OIL (UNII: ZBP1YXW0H8)
GRAPEFRUIT OIL (UNII: YR377U58W9)
FRAGRANCE GREEN APPLE ORC2001072 (UNII: U9GH30P956)
ISOPROPYL ALCOHOL (UNII: ND2M416302)
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)
COCO-CAPRYLATE/CAPRATE (UNII: 8D9H4QU99H)
ORANGE OIL (UNII: AKN3KSD11B)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77857-832-01	51 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/17/2021	
2	NDC:77857-832-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/17/2021	
3	NDC:77857-832-32	946 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/17/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/17/2021	

Labeler - Mega Creation Inc. (014096208)

Registrant - Mega Creation Inc. (014096208)

Establishment

Name	Address	ID/FEI	Business Operations
Mega Creation Inc.		014096208	manufacture(77857-832)

Revised: 11/2021

Mega Creation Inc.