SKINTILLATE HAND SANITIZER- alcohol, isopropyl alcohol solution
D-Time Limited Liability Company

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.

Active Ingredients

Alcohol (80%)
Ethyl Alcohol (75%), Isopropyl Alcohol (5%)

Purpose

Antiseptic skin cleanser

Use

For personal hand hygiene to help prevent the spread of bacteria.

Warnings

For external use only.

When using this product avoid contact with eyes. If contact occurs, rinse thoroughly with water. Stop use and consult a healthcare professional if irritation develops.

Flammable. Keep away from heat and flame.

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

Directions

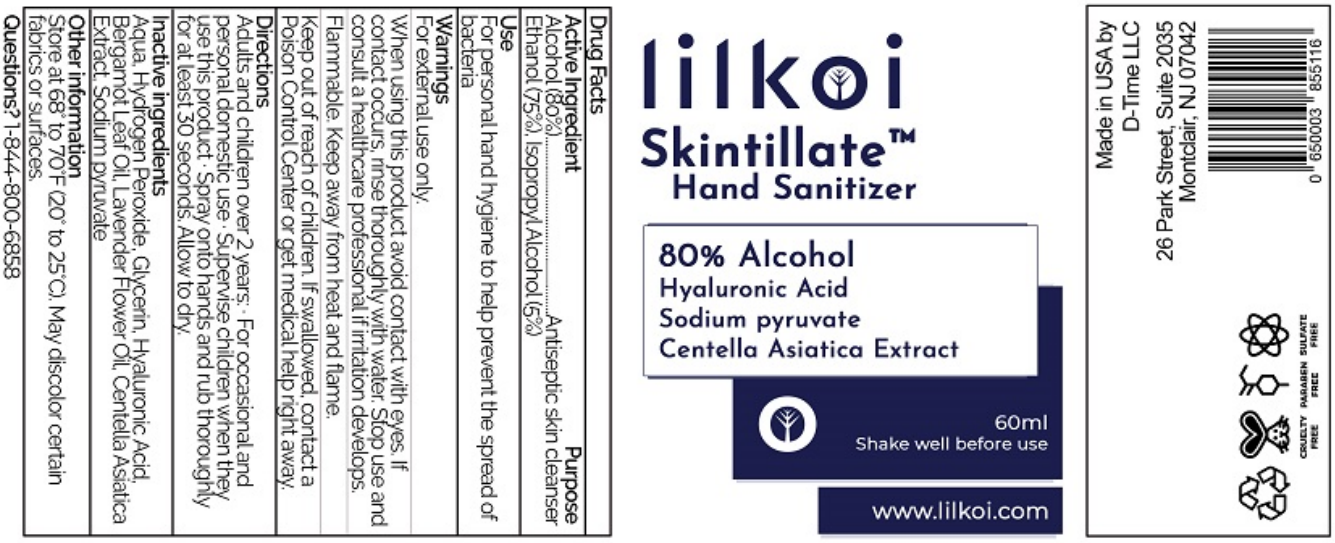
Adults and children over 2 years: For occasional and personal domestic use. Supervise children when they use this product. Spray onto hands and rub thoroughly for at least 30 seconds. Allow to dry.

Inactive ingredients

Water, Hydrogen Peroxide , Glycerin, Hyaluronic Acid, Bergamot leaf oil, Lavender Flower Oil, Centella Asiatica Extract, Sodium pyruvate

Other information
Store at 68^0 to 70^0 F (20^0 to 25^0C). May discolor certain fabrics or surfaces.
Questions? 1-844-800-6858

Package Label - Principal Display Panel



SKINTILLATE HAND SANITIZER

alcohol, isopropyl alcohol solution

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75306-003
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
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	5 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYALURONIC ACID (UNII: S270N0TRQY)	
BERGAMOT OIL (UNII: 39W1PKE3JI)	
LAVENDER OIL (UNII: ZBP1YXW0H8)	
CENTELLA ASIATICA (UNII: 7M867G6T1U)	
SODIUM PYRUVATE (UNII: POD38AIF08)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date

1	NDC:75306-003-01	30 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/01/2020	
2	NDC:75306-003-02	50 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/01/2020	
3	NDC:75306-003-03	60 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/01/2020	
4	NDC:75306-003-04	100 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/01/2020	
5	NDC:75306-003-05	120 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/01/2020	
6	NDC:75306-003-06	160 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/01/2020	
7	NDC:75306-003-07	200 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/01/2020	
8	NDC:75306-003-08	250 mL in 1 BOTTLE, PLASTIC; 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	part333A	05/01/2020	

Labeler - D-Time Limited Liability Company (081728006)

Establishment

Name	Address	ID/FEI	Business Operations
D-Time Limited Liability Company		081728006	manufacture(75306-003)

Revised: 5/2020 D-Time Limited Liability Company