LIFESHIELD HAND SANITIZER 70% ETOH- ethyl alcohol gel LifeTech Pharma LLC.

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.

Hand Sanitizer

This hand Sanitizer was manufactured according to the Federal Register 21 CFR 310, utilizing:

- a. USP/FCC grade Ethyl alcohol 99%
- b. Distilled/purified water
- c. Hydroxypropylcellulose (as Vegetable Cellulose)
- d. Glycerin USP/FCC grade
- e. Limonene (as fragrance)
- f. Aloe Vera extract

Active Ingredient(s)

Ethyl alcohol 70%

Purpose

Antiseptic, Hand Sanitizer

Uses

Decreases bacteria on skin.

Warnings

For external use only. Flammable, keep away from flames.

When using this product

When using this product keep out of eyes. In case of contact, flush eyes with water.

Stop use

Stop use and ask doctor if irritation and redness develop.

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

Directions

- Rub hands thoroughly with product and allow to dry.
- For children under 6, use only under adult supervision.

Other information

- Store below 110°F (43°C).
- May discolor certain fabrics or surfaces.

Inactive ingredients

Inactive ingredients Distilled Water, Structured Silver, Vegetable Cellulose, Glycerin, Aloe Vera, Fragrance.

Questions

Questions? Call 1-385-244-0470 Monday through Friday 8:30 AM to 5:00 PM MST.

2 oz Packaging & Label - Principal Display Panel

2 FL OZ (59.15 mL) NDC: 75270-104-02





128 oz Packaging & Label - Principal Display Panel

1 Gallon (3.78 L) 128 FL Oz - NDC: 75270-104-28



1 GAL (3.78L) 128 OZ





8 oz Packaging & Label - Principal Display Panel

8 FL Oz (236 mL) - NDC: 75270-104-08



16 oz Packaging & Label - Principal Display Panel

16 FL Oz (473 mL) - NDC: 75270-104-16



32 oz Packaging & Label - Principal Display Panel

32 FL Oz (946 mL) - NDC: 75270-104-32



LIFESHIELD HAND SANITIZER 70% ETOH ethyl alcohol gel Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:75270-104

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TOPICAL

Active ingredient/Active triviety			
Ingredient Name	Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70.7 mL in 100 mL	

Inactive Ingredients				
Ingredient Name	Strength			
LIMO NENE, (+)- (UNII: GFD7C86Q1W)	0.1 mL in 100 mL			
ALOE VERA LEAF POLYSACCHARIDES (UNII: W21O437517)	0.01 mL in 100 mL			
GLYCERIN (UNII: PDC6A3C0OX)	0.7 mL in 100 mL			
WATER (UNII: 059QF0KO0R)	28.5 mL in 100 mL			
HYDRO XYPRO PYL CELLULO SE, UNSPECIFIED (UNII: 9 XZ8 H6 N6 OH)	0.01 mL in 100 mL			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75270-104- 02	59.15 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	04/28/2020	
2	NDC:75270-104- 28	3780 mL in 1 JUG; Type 0: Not a Combination Product	04/28/2020	
3	NDC:75270-104- 08	236 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	04/28/2020	
4	NDC:75270-104- 32	946 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	04/28/2020	
5	NDC:75270-104- 16	473 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	04/28/2020	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	04/28/2020		

Labeler - LifeTech Pharma LLC. (057492812)

Registrant - LifeTech Pharma LLC. (057492812)

Establishment				
Name	Address	ID/FEI	Business Operations	
LifeTech Pharma		057492812	manufacture(75270-104)	

Revised: 5/2020 LifeTech Pharma LLC.