LXR HEALTHCARE- hand sanitizer spray liquid LXR Biotech 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.

LXR-Sanitizer Spray Blue-Label 86% Alcohol

Active Ingredient(s)

Alcohol 86% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer

Use

Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

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

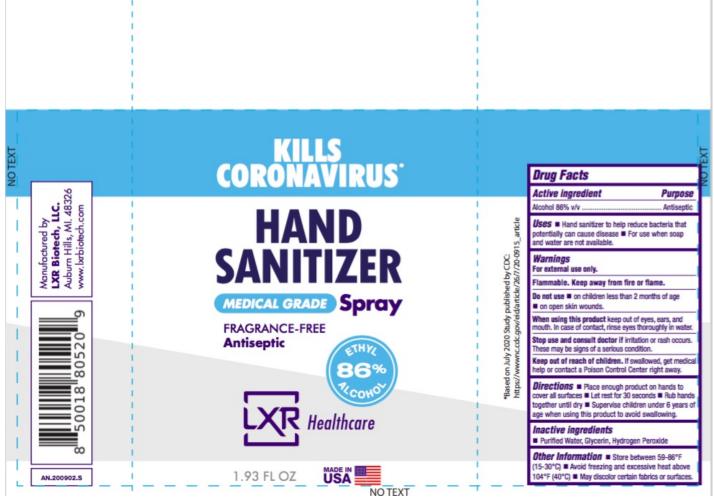
- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

Store between 59-86F (15-30C) Avoid freezing and excessive heat above 104F (40C) May discolor certain fabrics or surfaces

Inactive ingredients

Package Label - Principal Display Panel



57 mL NDC: 78299-131-11

LXR HEALTHCARE					
hand sanitizer spray liquid					
Product Information					
Product T ype	HUMAN OTC DRUG	Item Code (Source)		NDC:78299-131	
Route of Administration	TOPICAL				
Active Ingredient/Active Moi	ety				
Ingredient Name			Basis of Strength	n Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)			ALCOHOL	86 mL in 100 mL	
Inactive Ingredients					
Ingredient Name			Strength		
GLYCERIN (UNII: PDC6A3C0OX)					
HYDROGEN PEROXIDE (UNII: BBX06	50 AN9 V)				

WATER (UNII: 059QF							
Packaging							
# Item Code	Package Description	Marketing Start Date	Marketing End Date				
1 NDC:78299-131-11	57 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/12/2020					
Marketing Information							
Marketing Catego	ry Application Number or Monograph Citation	Marketing Start Date	Marketing End Date				
OTC monograph not f	inal part333A	09/12/2020					

Labeler - LXR Biotech LLC (117520926)

Establishment					
Name	Address	ID/FEI	Business Operations		
LXR Biotech LLC		117520926	manufacture(78299-131)		

Revised: 9/2020

LXR Biotech LLC