LINSTOL REFRESHING AND CLEANSING HAND SANITIZER- alcohol gel
Linstol USA, 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.

Linstol Refreshing & Cleaning Hand Sanitizer

Drug Facts

Active ingredient

Ethyl Alcohol 75%

Purpose

Antiseptic

Uses

Hand sanitizer to help reduce bacteria on the skin

Warnings

For external use only

Flammable. Keep away from fire or flame.

When using this product do not use in or near the eyes. If contact occurs, rinse eyes thoroughly with water.

Stop use or ask a doctor if irritation or rash appears and lasts.
- You may report side effects to FDA at **1-800-FDA-1088**.

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

Directions
- Wet hands thoroughly with product and rub hands together briskly until dry.
- Children under 6 years of age should be supervised when using this product.

Inactive ingredients

Water (Aqua), Carbomer, Fragrance, Glycerin, Racementhol, Trolamine.

Distributed by: Linstol USA, LLC
Naples, FL 34114

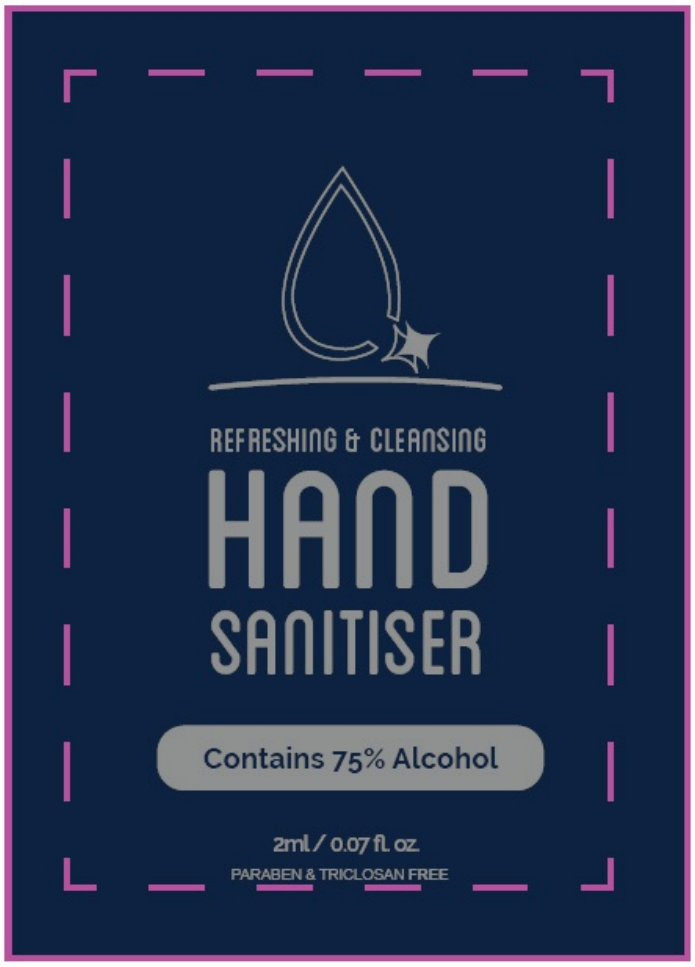
PRINCIPAL DISPLAY PANEL - 2 ml Packet

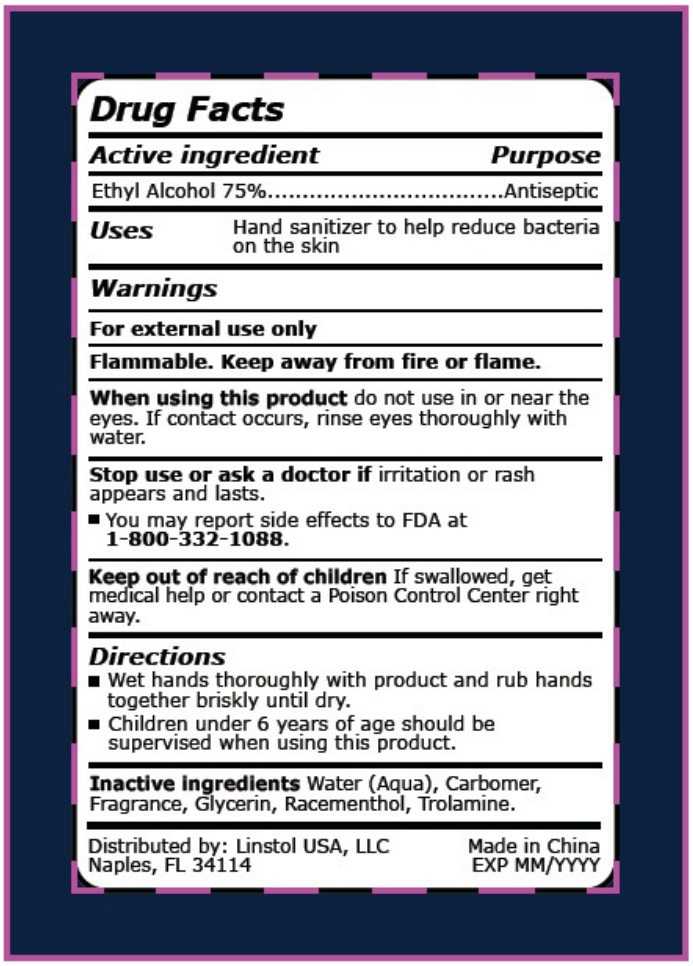
REFRESHING & CLEANSING

HAND
SANITISER

Contains 75% Alcohol

2ml / 0.07 fl. oz.
PARABEN & TRICLOSAN FREE





LINSTOL REFRESHING AND CLEANSING HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:76982-3864
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

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
RACEMENTHOL (UNII: YS08XHA860)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0KO0R)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76982-3864-0	2 mL in 1 PACKET; Type 0: Not a Combination Product	05/20/2020	
2	NDC:76982-3864-1	15 mL in 1 TUBE; Type 0: Not a Combination Product	05/20/2020	
3	NDC:76982-3864-2	30 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/20/2020	
4	NDC:76982-3864-3	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/20/2020	
5	NDC:76982-3864-4	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/20/2020	
6	NDC:76982-3864-5	236 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/20/2020	
7	NDC:76982-3864-6	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/20/2020	
8	NDC:76982-3864-7	532 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/20/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/20/2020	

Labeler - Linstol USA, LLC (828039995)

Revised: 12/2022 Linstol USA, LLC