NATURALIXIR- alcohol gel
North Atlantic Chemicals and Technologies 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.

NaturAlixir

Drug facts

Active ingredient(s)

Alcohol 70% w/w

Purpose

Antiseptic

Use(s)

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 eye, rinse eyes thoroughly with water.

Stop use and see 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.

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

Inactive ingredients Distilled Water, All-Natural Fragrance Oil, Polyacrylate crosspolymer-6, Aloe Vera Leaf Juice, Vegetable Glycerin.

With Natural Ingredients

No Parabens, Aldehydes

No Animal-based Ingredients

Vegan

Packaging

NATURALIXIR

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73835-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	

AMMONIUM ACRYLOYLDIMETHYLTAURATE, DIMETHYLACRYLAMIDE, LAURYL METHACRYLATE AND LAURETH-4 METHACRYLATE COPOLYMER, TRIMETHYLOLPROPANE TRIACRYLATE CROSSLINKED (45000 MPA.S) (UNII: Q7UI015FF9)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73835-001-02	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
2	NDC:73835-001-04	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
3	NDC:73835-001-08	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
4	NDC:73835-001-32	946 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
5	NDC:73835-001-64	1893 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
6	NDC:73835-001-10	3785 mL in 1 CONTAINER; Type 0: Not a Combination Product	03/30/2020	
7	NDC:73835-001-25	9463 mL in 1 CONTAINER; Type 0: Not a Combination Product	03/30/2020	
8	NDC:73835-001-50	18927 mL in 1 CONTAINER; Type 0: Not a Combination Product	03/30/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/30/2020	

Labeler - North Atlantic Chemicals and Technologies LLC (081316806)

Establishment

Name	Address	ID/FEI	Business Operations
North Atlantic Chemicals and Technologies LLC		081316806	manufacture(73835-001)

Revised: 4/2020 North Atlantic Chemicals and Technologies LLC