

NATURE DROPS HAND SANITIZER- alcohol liquid**Bioplasticos y Polimeros BIOFASE, S.A.P.I. de C.V.**

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.

Nature Drops Hand Sanitizer**DRUG FACT CHART****Active Ingredients:**

Ethyl Alcohol at 70%

Purpose:

Antiseptic

Uses:

Hand Sanitizer to help reduce germs. For use when soap and water not available.

Warnings:

For external use only, Flammable. Keep away from open flame and sources of heat or flame.

Do not use:

In children less than 2 months of age. Open skinwounds.

When using this product:

Keep out of eyes, ears, and mouth. In case of contact with eyes, rinse immediately with plenty of water and seek medical advice.

Stop use and ask doctor:

If an irritation or rash occurs.

Keep out of reach of children:

If swallowed, seek medical advice or contact a poison control center immediately.

Directions:

Apply a small amount of product into hands and rub between both side of hands until dry. Supervise children under 6 years of age when using this product to avoid swallowing.

Other information:

Store below 110°F (43°C). Avoid freezing. May discolor certain fabrics or surface.

Inactive Ingredients:

Water, Glycerin, Hydroxyethyl Cellulose, fragrance.

Package Labeling:



NatureDrops®

Products from the Earth

HAND SANITIZER

100% PLANT BASED

(Ethyl Alcohol Anticeptic 70%)
Non Sterile and Topical Solution



Ethyl Alcohol made from plant sugars.



Thickener agent made from cotton.



Moisturizer made from natural oils.



Plant based fragrance.

1000 ml (33.8 fl. oz.)

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CONTACT US

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Bottled and Packed by:
Bioplasticos y Polimeros Biofase
SAPI de CV Circuito Parque Industrial 159
Consuelo Vázquez, Morelia Mich.
México ZC 58200

NATURE DROPS HAND SANITIZER

alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77452-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)			ALCOHOL	0.7 mL in 1 mL
Inactive Ingredients				
Ingredient Name				Strength
WATER (UNII: 059QF0KO0R)				
GLYCERIN (UNII: PDC6A3C0OX)				
HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77452-001-01	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/24/2020	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final		part333E	04/24/2020	

Labeler - Bioplasticos y Polimeros BIOFASE, S.A.P.I. de C.V. (951577065)

Revised: 5/2020

Bioplasticos y Polimeros BIOFASE, S.A.P.I. de C.V.