NONGBIAE HAND SANITIZER- ethyl alcohol gel Evergreenplus

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

82390-102 NONGBIAE Hand Sanitizer

Active Ingredient

Ethyl Alcohol 60%

Purpose

Antiseptic

Uses

Hand sanitizer to decrease bacteria on the skin that could cause disease.

Warnings

Keep away from heat and flame. For external use only. Flammable.

Do not use

Stop using it if you have severe red spots or itching.

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

Stop use

Stop use and ask a doctor if irritation or rash occurs.

When using

- ·Keep out of eyes. In case of contact with eyes, flush thoroughly with water. ·Avoid contact with broken skin.
- ·Do not inhale or ingest.

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.

Inactive Ingredients

Water, Citric acid, Protein, Fatty acid ester, Glycerin

Label

NONGBIAE Hand Sanitizer Gel

[Drug Facts]

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Packing Label 100ml (3.38oz)

Manufacturing number and expiration date

Separate indication

Manufacturer and Manufacturer Seller

Evergreenplus

112-8, Jiji-gil, Beomseo-eup, Ulju-gun, Ulsan, 44918

Republic of Korea

NONGBIAE HAND SANITIZER

ethyl alcohol gel

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82390-102	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	60 mL in 100 mL	

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)		
GLYCERIN (UNII: PDC6A3C0OX)		
HYDROLYZED ORYZANOL (UNII: 30E0DD115J)		
ISOPROPYL BEHENATE (UNII: 1BWO0L9BHI)		

Packaging					
	# Item Code Package Description		Marketing Start Date	Marketing End Date	
	1	NDC:82390- 102-01	100 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/16/2021	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved drug		11/16/2021		

other 11/10/2021

Labeler - Evergreenplus (695376425)

Establishment				
Name	Address	ID/FEI	Business Operations	
Evergreenplus		695376425	manufacture(82390-102)	

Revised: 11/2021 Evergreenplus