RAIN FOREST HAND SANITIZER- alcohol gel
Antrix Distribution 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.

Rain Forest Hand Sanitizer - Antrix

Drug Facts

Active ingredients

Alcohol 70% v/v

Purpose

Antiseptic

Uses

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

- On 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 thoroughly with water.

Stop use and ask a doctor if

- Stop use and seek 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 15-30C (59-86F).
- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients

Water, Carbomer, Glycerin, Triethanolamine, Iodopropynyl Butylcarbamate, Methylisothiazolinone.

Questions?

call 888 919 9223

Company Information

DISTRIBUTED BY
Antrix Distribution,
2013 Murcott drive suite E
St. Cloud Fl 34771
Made in Guatemala

Product Packaging - 240 ml

RAIN
FOREST
HAND
SANITIZER
NON STERILE SOLUTION
with moisturizing ingredients
ALCOHOL 70%
BY
NAKED NATURALS
8 fl. oz. / 240ml

RAIN FOREST HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:77646-001
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
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0KO0R)	

Packaging

			Marketing Start	Marketing End

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77646-001-01	60 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/12/2020	
2	NDC:77646-001-02	120 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/12/2020	
3	NDC:77646-001-03	240 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/12/2020	
4	NDC:77646-001-04	3780 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/12/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/12/2020	

Labeler - Antrix Distribution LLC (081436980)

Revised: 5/2020 Antrix Distribution LLC