NIFOLA HAND SANITIZER WITH ALOE VERA AND VITAMIN E- alcohol gel
Yiwu Yangjie Daily Chemicals Co.,Ltd.

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.

Drug facts

Active ingredient Purpose

Ethyl Alcohol 70% V/V ... Antiseptic

- help decrease bacteria on the hands.
- recommended for repeated use.

if swallowed, get meical help or contact a poison control center right away.

Dispense a small amount on palms. Rub lightly until dry. Do not rinse.

For external use only.
Flammable. Keep away from fire or flame.

Avoid contact with eyes, in case of contact, rinse with water immediately.

If irritation develops, discontinue use and consult a doctor.

OTHER INFORMATION: store under 105□.

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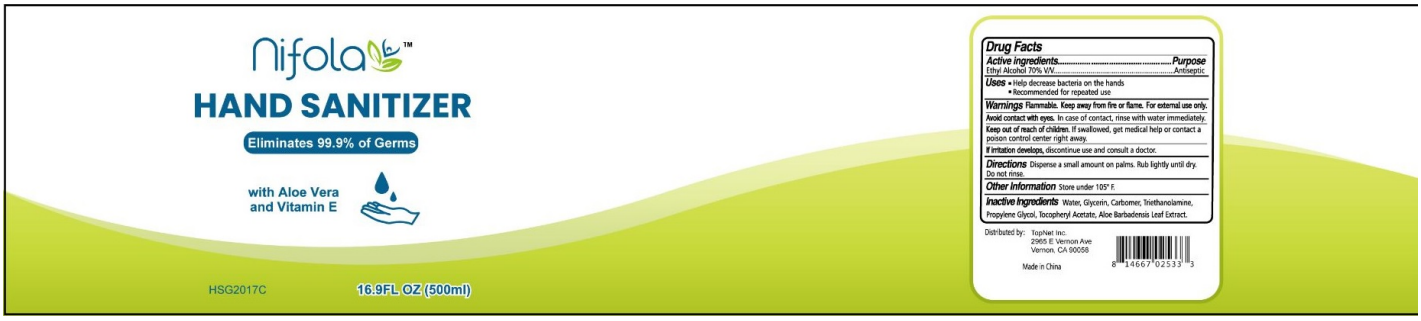
Water, Glycerin, Carbomer, Triethanolamine, Propylene Glycol, Tocopheryl Acetate, Aloe Barbadensis Leaf Extract



NIFOLA HAND SANITIZER WITH ALOE VERA AND VITAMIN E

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:74149-022
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)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
TROLAMINE (UNII: 9O3K93S3TK)	
ALPHA-TOCOPHEROL (UNII: H4N855PNZ1)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74149-022-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/12/2020	
2	NDC:74149-022-02	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/12/2020	
3	NDC:74149-022-03	53 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/12/2020	
4	NDC:74149-022-04	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/12/2020	
5	NDC:74149-022-05	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/12/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/12/2020	

Labeler - Yiwu Yangjie Daily Chemicals Co.,Ltd. (529648827)

Establishment

Name	Address	ID/FEI	Business Operations
Yiwu Yangjie Daily Chemicals Co.,Ltd.		529648827	manufacture(74149-022)

Revised: 6/2020 Yiwu Yangjie Daily Chemicals Co.,Ltd.