NEXSENSE LIQUID HAND SANITIZER MOISTURIZING WITH ALOE- alcohol solution Taizhou Yadai Daily-Use 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.

52989-003

Active Ingredient(s)

Ethyl Alcohol 62% v/v

Purpose

Antiseptic

Use

Hand sanitizer to help decrease bacteria on the skin

Recommended for repeated use

Warnings

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

Do not apply around eyes or use in ears or mouth

Stop use and contact a doctor if redness or irritation develop or persist for more than 72 hours

Keep out of reach of children. If swallowed, get medical help or call a Poison Control Center right away.

Directions

- Pump as needed into your palms and thoroughly spead on both hands
- rub into skin until dry

Other information

- Store at room temperature no higher than 45C (115F) and under shade
- Keep in closed containers and free from direct sunlight

Inactive ingredients

Water, Glycerol, Aminomethyl propanol, Aloe Vera gel, Vitamin E, Carbomer, Frangrance





Strength

NEXSENSE LIQUID HAND SANITIZER MOISTURIZING WITH ALOE

alcohol solution

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:52989-003

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name
Basis of Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)
ALCOHOL (UNII: 3K9958V90M) ALCOHOL 62 mL in 100 mL

Inactive Ingredients

CARBOMER 934 (UNII: Z135WT9208)

Ingredient Name

AMINO METHYLPRO PANO L (UNII: LU49 E6626Q)

ALOE VERA WHOLE (UNII: KIZ4X2EHYX)

GLYCERIN (UNII: PDC6A3C0OX)

TROLAMINE (UNII: 903K93S3TK)

.ALPHA.-TO COPHERO L ACETATE (UNII: 9E8 X80 D2L0)

Packaging	
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#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52989-003-01	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/19/2020	
2	NDC:52989-003-02	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/19/2020	

3 NDC:52989-003-03 500							
Marketing Information							
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date				
OTC monograph not final	part333A	06/19/2020					

Labeler - Taizhou Yadai Daily-Use Co., Ltd. (529899639)

Establishment							
Name	Address	ID/FEI	Business Operations				
Taizhou Yadai Daily-Use Co., Ltd.		529899639	manufacture(52989-003)				

Revised: 6/2020 Taizhou Yadai Daily-Use Co., Ltd.