AURAFRESH INSTANT HAND SANITIZER VITAMIN E- alcohol gel Ningbo Pulisi Daily Chemical Products 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% ... Antiseptic

Hand sanitizer to help decrease bacteria on the skin.

If swallowed, get medical help or contact local poison control center right away.

Children under 6 years of age should be supervised when using this product.

keep out of eyes

when water, soap and towel are not available

For external use only.

Flammable. Keep away from hear or flame.

When using this products, do not use in or the eyes. In case of contact, rinse eyes thoroughly with water.

Stop using and ask a doctor, if irritation or rash appears and lasts.

place enough product in your palms to thoroughly spread on both hands, and rub into the skin until dry.

Water, Carbomer, Glycerin, Propylene Glycol, Triethanolamine, acrylates/c10-30 alkyl acrylate crosspolymer, isopropyl alcohol, aminomethyl propanol, isopropyl mysistate, caprylyl glycol, phenoxyethanol, Tocopheryl Acetate, Fragrance





AURAFRESH INSTANT HAND SANITIZER VITAMIN E

alcohol gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:40104-284

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name
Basis of 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: ZY8 1Z8 3H0 X)

CARBOMER 934 (UNII: Z135WT9208)

GLYCERIN (UNII: PDC6A3C0OX)

.ALPHA.-TO COPHEROL ACETATE, D- (UNII: A7E6112E4N)

TROLAMINE (UNII: 903K93S3TK)

CARBOMER COPOLYMER TYPE A (UNII: 71DD5V995L)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
CAPRYLYL GLYCOL (UNII: 00 YIU5438 U)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

Packaging						
7	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:40104-284-01	2 in 1 PACKAGE	08/10/2017			
1		60 mL in 1 BOTTLE; Type 0: Not a Combination Product				

Marketing Information							
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date				
OTC monograph not final	part333A	08/10/2017					

Labeler - Ningbo Pulisi Daily Chemical Products Co., Ltd. (529047265)

Registrant - Ningbo Pulisi Daily Chemical Products Co., Ltd. (529047265)

Establishment						
Name	Address	ID/FEI	Business Operations			
Ningbo Pulisi Daily Chemical Products Co., Ltd.		529047265	manufacture (40 10 4-28 4)			

Revised: 8/2017 Ningbo Pulisi Daily Chemical Products Co., Ltd.