INSTANT HAND SANITIZER- instant hand sanitizer gel ADF HI-TECH DISINFECTANTS 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.

Keep in dry and cool places, away from fire.

carbomer[]glycerine[]purified water

Put proper amount of the product on palm and rub thoroughly until dry.

Ethanol

keep out of reach of children

Disinfection Sterilization No Rinseing

Flammable. For external use only. Avoid contact with eyes and broken skin. Discontinue use if irritation develops.

Drug Facts

Active ingredient

Ethanol alcohol 75%(v/v).....Antiseptic

Purpose

Inactive ingredients

Purified water, carbomer, glycerine

Uses

Help reduce pathogens on hand, skin and objects surfaces.

Storage

Keep in dry and cool places, away from fire and out of children's reach. Valid for 24 months.

Directions

Put proper amount of the product on palm and rub thoroughly until dry.

Warnings

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Flammable. For external use only. Avoid contact with eyes and broken skin. Discontinue use if irritation develops.



Instant Hand Sanitizer

Kills **99.99%** of many common harmful germs

17FL 0Z(502ml)

INSTANT HAND SANITIZER instant hand sanitizer gel **Product Information** NDC:52923-002 **Product** Type HUMAN OTC DRUG Item Code (Source) **Route of Administration** EXTRACORPOREAL **Active Ingredient/Active Moiety Ingredient Name Basis of Strength** Strength ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII: 3K9958V90M) ALCOHOL 351.4 mL in 502 mL **Inactive Ingredients Ingredient Name** Strength

GLYCERIN (UNII: PDC6A3C0OX)										
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)										
W	WATER (UNII: 059QF0KO0R)									
Product Characteristics										
Color			S	Score						
Shape			S	Size						
Flavor			Ŀ	Imprint Code			KONGPILI			
Contains										
Packaging										
#	Item Code	Р	Package Description		Marketing St	art Date	Marketing End Date			
1	NDC:52923-002-01	502 mL in 1 BOTT	2 mL in 1 BOTTLE; Type 0: Not a Combination Product			05/19/2020				
Marketing Information										
	Marketing Categor	y Applicatio	on Num	iber or Monograph Citation	Marketing St	art Date	Marketing End Date			
OTC monograph not final part333E					05/19/2020					

Labeler - ADF HI-TECH DISINFECTANTS CO., LTD. (529220014)

Establishment

Name	Address	ID/FEI	Business Operations
ADF HI-TECH DISINFECTANTS CO., LTD.		529220014	manufacture(52923-002)

Revised: 5/2020

ADF HI-TECH DISINFECT ANTS CO., LTD.