

PURE CHOICE INSTANT HAND SANITIZER- alcohol gel

Shanghai Medical Instruments 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.

Pure Choice Instant Hand Sanitizer

Drug Facts

Active ingredient

Ethyl alcohol 75% v/v

Purpose

Antimicrobial

Use:

Hand sanitizer to help reduce bacteria on the skin

Warnings

Flammable. Keep away from fire or flame

For external use only

When using this product

do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water

Stop use and ask a doctor if

irritation or rash appears and lasts

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions:

1)Put enough product in your palm to cover hands and rub hands together briskly until dry. 2)

Children under 6 years of age should be supervised when using

Inactive ingredients

Glycerin, Water, Carbomer, aloe extraction, Triethanolamine, Ammonium Lauryl Sulfate, Hyaluronic Acid, Aloe Extract, Parfum

Question? 1-866-222-3181.

Package Labeling:

VALUEPACK EXTRA 10%

PureChoice

INSTANT
HAND
SANITIZER

ADVANCED FORMULA
WITH MOISTURIZERS

REFRESHING ALOE

8.8 FL OZ (260ml)

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PURE CHOICE INSTANT HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74854-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.75 mL in 1 mL

Inactive Ingredients

Ingredient Name				Strength
GLYCERIN (UNII: PDC6A3C0OX)				
WATER (UNII: 059QF0KO0R)				
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)				
TROLAMINE (UNII: 9O3K93S3TK)				
AMMONIUM LAURYL SULFATE (UNII: Q7AO2R1M0B)				
HYALURONIC ACID (UNII: S270N0TRQY)				
ALOE (UNII: V5VD430YW9)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74854-001-26	260 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/22/2020	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final		part333E	04/22/2020	

Labeler - Shanghai Medical Instruments Co.,Ltd (545004038)

Establishment			
Name	Address	ID/FEI	Business Operations
Shanghai Medical Instruments Co.,Ltd		545004038	manufacture(74854-001)

Revised: 4/2020

Shanghai Medical Instruments Co.,Ltd