BENEMAX ADVANCED HAND SANITIZER- isopropyl alcohol liquid Nugale Pharmaceutical Inc

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

Isopropyl alcohol 70%, USP

Purpose

Antiseptic

Warnings

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

Do not use

- on children less than 2 months of age
- on open skin wounds

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

Directions

• Place enough product on hands or affected area to cover all surfaces. Rub hands together until dry.

• Supervise children under 6 years of age when using this product to avoid swallowing.

Uses

Antiseptic and Hand Sanitizer to help reduce bacteria that may cause disease. For use when soap & water are unavailable.

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away at 1-800-222-1222.

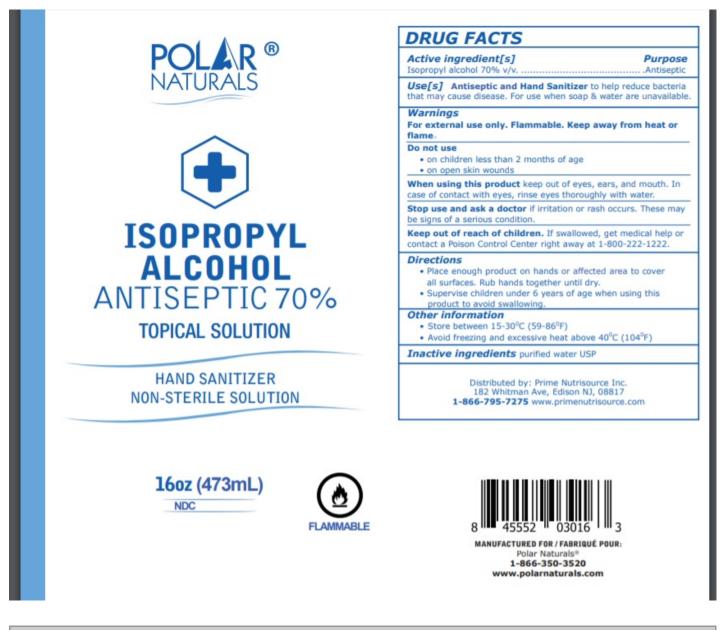
Inactive ingredient

Water

Other information

Store between 15 °C and 30 °C. Avoid freezing and excessive heat above 400 °C (104 °F)

Product label



BENEMAX ADVANCED HAND SANITIZER

isopropyl alcohol liquid

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:73700-004

Route of Administration TOPICAL

Ac	Active Ingredient/Active Moiety								
		In	gredient Name		Basis of Strength	Strongth			
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOUNII: ND2M416302)			ND2M416302) (ISOPROPYL ALCOHOL -		ISOPROPYL ALCOHOL	70 mL in 100 mL			
Inactive Ingredients									
Ingredient Name WATER (UNII: 059QF0KO0R)					Strength				
Packaging									
Pa	ackaging			•••					
Pa #	ackaging Item Code		Package Description	Mar	keting Start Date	Marketing Enc Date			
#	Item Code	473 mL in 1 Combinatior	BOTTLE, PLASTIC; Type 0: Not a	Mar 02/25,	Date	-			
#	Item Code		BOTTLE, PLASTIC; Type 0: Not a		Date	-			
#	Item Code	Combinatior	BOTTLE, PLASTIC; Type 0: Not a Product		Date	-			
#	Item Code NDC:73700- 004-01	Combination	BOTTLE, PLASTIC; Type 0: Not a Product	02/25,	Date	-			

Labeler - Nugale Pharmaceutical Inc (202595872)

Establishment								
Name	Address	ID/FEI	Business Operations					
Nugale Pharmaceutical Inc		202595872	manufacture(73700-004)					

Revised: 3/2021

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Nugale Pharmaceutical Inc