

HYGEN HAND SANITIZER- isopropyl alcohol liquid

Lonza, LLC

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.

HYGEN® Hand Sanitizer

Drug Facts

Active Ingredient[s]

Isopropyl alcohol 75% v/v

Purpose

Antiseptic

Use(s)

Hand sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

Warnings

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

Do not use

- In children less than 2 months of age.
- On open skin wounds.

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

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

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

Directions

- Place enough product on hands to cover all surfaces. Rub hands together until dry .
- Supervise children under 6 years of age when using this product to avoid swallowing .

Other information

- Store between 15-30°C (59-86°F)
- Avoid freezing and excessive heat above 40°C (104°F)

Inactive ingredients

glycerin, hydrogen peroxide, purified water USP

Manufactured by Lonza, LLC

412 Mt. Kemble Avenue, Suite 2005,

Morristown, NJ 07960

PRINCIPAL DISPLAY PANEL - 3.79 L Container Label

Lonza

HYGEN® Hand Sanitizer

ISOPROPYL ALCOHOL ANTISEPTIC 75%
TOPICAL SOLUTION

HAND SANITIZER
NON-STERILE SOLUTION

NET CONTENTS: 1 Gal

HYGEN® Hand Sanitizer

PROD. No.

LOT No.

Net Weight:

UN1993

Flammable liquids, n.o.s.
(Isopropanol)



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LOT No.

Net Weight:

Manufactured by Lonza, LLC
412 Mt. Kemble Avenue, Suite 200S,
Morristown, NJ 07960
Made in the USA

Remark:

CONSULT SAFETY DATA SHEET.

Revision date: 00-00-0000 0 000000039929 /

HYGEN HAND SANITIZER			
isopropyl alcohol liquid			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69284-005
Route of Administration	CUTANEOUS		
Active Ingredient/Active Moiety			

Ingredient Name		Basis of Strength	Strength	
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)		ISOPROPYL ALCOHOL	750 mL in 1 L	
Inactive Ingredients				
Ingredient Name			Strength	
HYDROGEN PEROXIDE (UNII: BBX060AN9V)				
GLYCERIN (UNII: PDC6A3C0OX)				
WATER (UNII: 059QF0K00R)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69284-005-01	18.93 L in 1 PAIL; Type 0: Not a Combination Product	03/31/2020	
2	NDC:69284-005-02	208 L in 1 DRUM; Type 0: Not a Combination Product	03/31/2020	
3	NDC:69284-005-03	3.79 L in 1 CONTAINER; Type 0: Not a Combination Product	03/31/2020	
4	NDC:69284-005-04	0.5 L in 1 CONTAINER; Type 0: Not a Combination Product	05/01/2020	
5	NDC:69284-005-05	0.25 L in 1 CONTAINER; Type 0: Not a Combination Product	05/01/2020	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC MONOGRAPH NOT FINAL	part333E	03/31/2020		

Labeler - Lonza, LLC (001643170)

Revised: 4/2020

Lonza, LLC