

GEL HAND SANITIZER- alcohol gel 74793

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.

Gel Hand Sanitizer PQ-37 with Fragrance

Active Ingredient(s)

Alcohol 70% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer

Use

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, 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.

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

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

Inactive ingredients

Water, Glycerin, Polyquaterniun-37, hydrogen peroxide. Fragrance

Package Label - Principal Display Panel

250 mL NDC: 74793-0005-1

500 mL NDC: 74793-0005-2

Drug Facts	Purpose
Active Ingredient Ethyl Alcohol 70% v/v	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 • on children less than two 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.	

Drug Facts [Cont.]
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 Water, Glycerin, Polyquaternium-37, Hydrogen Peroxide, Fragrance

3780 mL NDC: 74793-0005-3

Drug Facts	Purpose
Active Ingredient Ethyl Alcohol 70% v/v	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 • on children less than two 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.	
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 Water, Glycerin, Polyquaternium-37, Hydrogen Peroxide, Fragrance	



500 mL NDC # 74793-0005-2

GEL HAND SANITIZER			
alcohol gel			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74793-0005
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)		ALCOHOL	70 mL in 100 mL
Inactive Ingredients			
Ingredient Name			Strength
EAST INDIAN LEMONGRASS OIL (UNII: UP0M8M3VZ W)			0.04 mL in 100 mL
GLYCERIN (UNII: PDC6A3C0OX)			1.45 mL in 100 mL

HYDROGEN PEROXIDE (UNII: BBX060AN9V)	0.125 mL in 100 mL
WATER (UNII: 059QF0KO0R)	
POLYQUATERNIUM-37 (10000 MPA.S) (UNII: 41QWS48DFN)	0.67 mL in 100 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74793-0005-1	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/06/2020	
2	NDC:74793-0005-2	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/06/2020	
3	NDC:74793-0005-3	3780 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/06/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/06/2020	

Labeler - 74793 (117054225)

Registrant - Streamline Polymers LLC (117054225)

Establishment

Name	Address	ID/FEI	Business Operations
Streamline Polymers LLC		117054225	manufacture(74793-0005)

Revised: 1/2022

74793