PFP POLYCLEAN HAND SANITIZER - GEL 62- gel hand sanitizer gel
Pfp Industries, LLC (a Global Enterprise)

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.

PFP POLYclean Hand Sanitizer - Gel 62

Drug Facts

Active ingredient [s]
Ethyl Alcohol 62% 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, 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 15-30C (59-86F)
•Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients Ammonium Polyacryloyldimethyl Taurate*, Fragrance, Glycerin, Hydrogen peroxide, Isopropyl

Alcohol, Sterile water, Ticacel*

*May or May Not Contain

Product Labeling

PfP INDUSTRIES **LOT Number :**

POLYClean Hand Sanitizer—GEL 62 **MADE IN TEXAS**

FLAMMABLE LIQUID

3

DOT: UN1170, II/III, Flammable Liquid,
n.o.s (contains ethanol) Percent Active

Ethyl Alcohol : 62% (v/v)

Before handling this material, read accompanied Safety Data Sheet for more detailed safety, health and environmental data.

Net Vol : 325 gal

Address : PFP Industries LLC

34011 Sunset lane Brookshire TX 77423
Ph: 281 371 2000

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res 8.22.20

PFP POLYCLEAN HAND SANITIZER - GEL 62

gel hand sanitizer gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:77034-006
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 L in 100 L

Inactive Ingredients	
Ingredient Name	**Strength**
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
WATER (UNII: 059QF0KO0R)	

Other Ingredients		
Ingredient Kind	**Ingredient Name**	**Quantity**
May contain	**AMMONIUM POLYACRYLOYLDIMETHYL TAURATE (55000 MPA.S)** (UNII: F01RIY4371)	
May contain	**METHYLCELLULOSE (25 MPA.S)** (UNII: BI55GG2WLI)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:77034-006-01	1217 L in 1 TANK; Type 0: Not a Combination Product	07/23/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/23/2020	

Labeler - Pfp Industries, LLC (a Global Enterprise) (080458439)

Revised: 8/2020 Pfp Industries, LLC (a Global Enterprise)