

PFP POLYCLEAN HAND SANITIZER - GEL 80- 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 80

Drug Facts

Active ingredient [s]

Ethyl Alcohol 80% 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 Glycerin, IPA, Hydrogen Peroxide, Sterile Water, Ticacel*, Ammonium Polyacryloyldimethyl Taurate*, Fragrance

*May or May Not Contain

Product Labeling

Pfp INDUSTRIES

LOT Number :

POLYClean Hand Sanitizer—GEL 80

MADE IN TEXAS

FLAMMABLE LIQUID

3

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

Ethyl Alcohol : 80% (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 80

gel hand sanitizer gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77034-007
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	80 L in 100 L

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	
WATER (UNII: 059QF0K00R)	

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:77034-007-01	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)