PHARMACYS PRESCRIPTION HAND SANITIZER 80Z- ethyl alcohol gel NINGBO LIYUAN DAILY CHEMICAL PRODUCTS CO., LTD.

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 ingredients

Ethyl Alcohol 62 percent

Purpose

Antiseptic

Uses

Helps reduce bacteria on the skin that could cause disease. Recommended for repeated use.

Warnings

For external use only.

Do not swallow.

Flammable. Keep away from fire or flame.

Do not apply around eyes. Do not use in ears & mouth.

When using this product, avoid contact with eyes. In case of contact, flush eyes with water.

Stop use and ask a doctor if

redness or irritation develops and persists for more than 72 hours.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

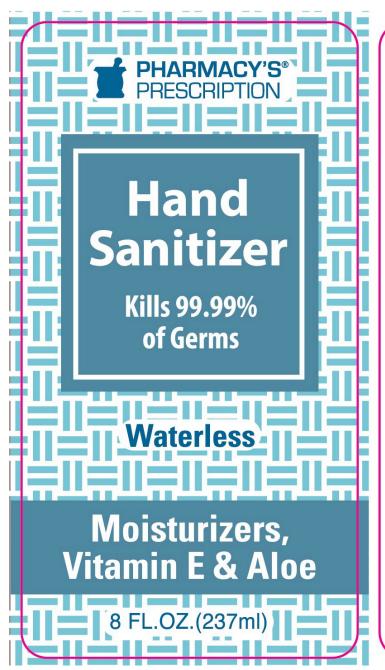
pump as needed into your palms and thoroughly spread on both hands, rub into skin until dry.

Other information

store at 20 Degree Celsius (68 to 77 Fahrenheit). may discolor fabrics.

Inactive ingredients

water, triethanolamine, glycerin, propylene glycol, tocopheryl acetate, aloe barbadensis gel, carbomber, fragrance.





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Distributed By:

American Consumer Products Corp. Vernon, CA 90058

NDC # 72197-011-08

Made in China

PHARMACYS PRESCRIPTION HAND SANITIZER 80Z

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76176-120
Poute of Administration	TOPICAL		

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

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)		
ALOE VERA LEAF (UNII: ZY81Z83H0X)		
CARBOMER 934 (UNII: Z135WT9208)		
TRIETHANOLAMINE BENZOATE (UNII: M3EN4GC19W)		
GLYCERIN (UNII: PDC6A3C0OX)		

l	Packaging					
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
		NDC:76176- 120-01	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/01/2018		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	10/01/2018		

Labeler - NINGBO LIYUAN DAILY CHEMICAL PRODUCTS CO., LTD. (530766098)

Registrant - NINGBO LIYUAN DAILY CHEMICAL PRODUCTS CO., LTD. (530766098)

Establishment				
Name	Address	ID/FEI	Business Operations	
NINGRO LIYUAN DAILY CHEMICAL PRODUCTS CO. LTD.		530766098	manufacture(76176-120)	

Revised: 10/2022 NINGBO LIYUAN DAILY CHEMICAL PRODUCTS CO., LTD.