PROTECTORX- alcohol gel PR Trading 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.

ProtectoRX

Drug Facts

Active ingredients

Ethyl alcohol 70%

Purpose

Antiseptic

Uses

- to decrease bacteria on the skin that could cause disease
- recommended for repeated use

Warnings

For external use only: hands Flammable, keep away from fire or flame.

When using this product

• keep out of eyes. In case of contact with eyes, flush thoroughly with water • avoid contact with broken skin • do not inhale or ingest

Stop use and ask a doctor if

• irritation and redness develop • condition 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

• wet hands thoroughly with product and allow to dry without wiping • for children under 6, use only under adult supervision • not recommended for infants

Other information

• do not store above 105°F • may discolor some fabrics • harmful to wood finishes and plastics

Inactive ingredients

Water, carbomer, aloe, glycerin, isopropyl myristate, tocopheryl acetate

Visit www.dibarlabs.com/Phone (281)-566-2682

Package Labeling:75408-002-01



Package Labeling:75408-002-02

Hand Sanitizer

Extra Strength





Disinfects and KILLS GERMS INSTANTLY

Contains moisturizing ingredients **Aloe & Vitamin E**

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Questions?

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Effective at eliminating common germs and bacteria. SDS-M0-15036 SDA-WI-2486 DSP-M0-28 DSP-M0-34



Imported by: **Dibar Labs, LLC.** Sugar Land, TX 77479

Distributed by: PR Trading LLC

42 Vondel Paseo del Parque San Juan, PR 00926

Cont. Net. 16 FL OZ (473.2 mL)

Made in México

PROTECTORX

alcohol gel

Product	Inform	nation
Product	INIAFE	narinn

Product Type HUMAN OTC DRUG Item Code (Source) NDC:75408-002

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.7 mL in 1 mL

Inactive Ingredients

Ingredient Name

Strength

WATER (UNII: 059QF0KO0R)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)	
ALOE (UNII: V5VD430 YW9)	
GLYCERIN (UNII: PDC6 A3C0 OX)	
ISOPROPYL MYRISTATE (UNII: 0 RE8 K4LNJS)	
.ALPHATO COPHEROL ACETATE (UNII: 9E8X80D2L0)	

	Packaging			
l	# Item Code	Package Description	Marketing Start Date	Marketing End Date
l	1 NDC:75408-002-01	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/26/2020	
	2 NDC:75408-002- 02	473.2 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/26/2020	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	04/26/2020		

Labeler - PR Trading LLC (117474793)

Revised: 4/2020 PR Trading LLC