PHARMAINT GEL HAND SANITIZER- ethyl alcohol 70% gel Laboratorio Pharma International Srl

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.

Pharmaint Gel Hand Sanitizer

Active Ingredient

Ethyl Alcohol 70%

Purpose

Antiseptic

Uses

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

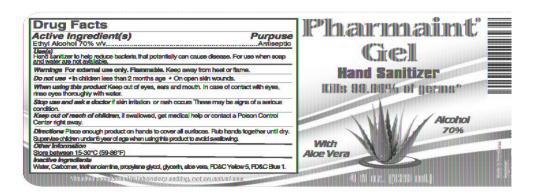
Store between 15-30C (59-86F).

Inactive Ingredients

Water, Carbomer, triethanolamine, propylene glycol, glycerine, aloe vera, FD&C Yellow 5, FD&C Blue 1

Pharmaint Gel Hand Sanitizer

ETIQUETA PARA FRASCO COSMÉTICO PLANO PET NATU



PHARMAINT GEL HAND SANITIZER

ethyl alcohol 70% gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:73182-101

Route of Administration TOPICAL

Active Ingredient/Active Moiety

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Ingredient Name	Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	84 mL in 120 mL	

nactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
ALOE VERA LEAF (UNII: ZY81Z83H0 X)		
POLYACRYLIC ACID (300000 MW) (UNII: A8371R0U5J)		
PROPYLENE GLYCOL ALGINATE (UNII: 26 CD3J2R0 C)		
TROLAMINE (UNII: 9O3K93S3TK)		
GLYCERIN (UNII: PDC6A3C0OX)		

Packaging					
ı	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
ı	1 NDC:73182-101-04	120 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020		

Marketing Inform	arketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part333A	05/01/2020			

Labeler - Laboratorio Pharma International Srl (816365775)

$\pmb{Registrant - \text{Laboratorio Pharma International Srl (816365775)}}$

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