BERNAL PREMIUM HAND SANITIZER- alcohol gel Inflatables 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.

bernal-PLD INFLATABLES LLC 70% Hand Sanitizer

Drug Facts

Active Ingredient(s)

Alcohol 70% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer

Use

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.

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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

Package Label - Principal Display Panel

220 mL NDC: NDC: 74541-570-08



BERNAL PREMIUM HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74541-570
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
l	Ingredient Name	Basis of Strength	Strength
l	ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients			
Ingredient Name	Strength		
CARBOMER 940 (UNII: 4Q93RCW27E)			
TROLAMINE (UNII: 9O3K93S3TK)			
WATER (UNII: 059QF0KO0R)			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74541-570- 01	30 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	03/30/2020	
2	NDC:74541-570- 02	60 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	03/30/2020	
3	NDC:74541-570- 04	130 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	03/30/2020	

4	NDC:74541-570- 08	220 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	03/30/2020
5	NDC:74541-570- 06	200 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	03/30/2020
6	NDC:74541-570- 17	500 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	03/30/2020
7	NDC:74541-570- 34	1000 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	03/30/2020
8	NDC:74541-570- 37	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/30/2020	

Labeler - Inflatables LLC (108619168)

Registrant - Margrey Industrial, S.A. de C.V. (814572392)

Establishment				
Name	Address	ID/FEI	Business Operations	
Margrey Industrial, S.A. de C.V.		814572392	manufacture(74541-570)	

Revised: 5/2020 Inflatables LLC